

**INHIBITORS OF CYCLIN DEPENDENT KINASES AS ANTI-CANCER  
AGENT**

**Related Applications**

This application is a continuation of PCT/GB02/01445, which was filed on March 26,  
5 2002, and which claims priority to GB 0107901.1, which was filed on March 29,  
2001. The entire contents of each of these applications are hereby incorporated herein  
by reference.

The present invention relates to 2-substituted 4-heteroaryl-pyrimidines, their  
preparation, pharmaceutical compositions containing them, and their use in the  
10 treatment of proliferative disorders such as cancer, leukemia, psoriasis and the like.

**Introduction and Summary of the Prior art**

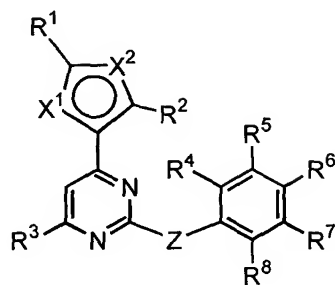
Certain 4,5,6-substituted-N-(substituted-phenyl)-2-pyrimidineamines having anti-  
asthmatic properties are disclosed in EP-A-233,461. Certain 4-heteroaryl-N-(3-  
15 substituted-phenyl)-2-pyridineamines possessing anti-proliferative properties and  
inhibiting protein kinases C, epidermal growth factor receptor-associated tyrosine  
protein kinase (EGF-R-TPK), as well as CDK1/cyclin B have been disclosed in  
WO95/09847 wherein the exemplified heteroaryl are pyridyl and indolyl.

J. Med. Chem. (1993) Vol. 36, pages 2716-2725, Paul, R. et al: discloses a further  
20 class of phenyl amino-pyrimidines possessing anti-inflammatory activity. These  
compounds include unsubstituted pyrrol groups, mono-substituted 2-thienyl groups  
and dimethyl-3-furyl groups at the 4-position of the pyrimidine ring.

It is an aim of the present invention to provide a further class of N-phenyl-2-  
pyrimidine anti-proliferative compounds. The compounds of the present invention  
25 have surprisingly been found to not to be inhibitors of protein kinase C. As discussed  
hereinafter, their activity may be demonstrated by inhibition of cell proliferation in  
cell lines and/or inhibition of cyclin dependent kinase enzymes.

**Summary of the Invention**

The first aspect of the present invention relates to compounds of general formula I:



**I**

wherein:

one of X<sup>1</sup> and X<sup>2</sup> is NR<sup>10</sup> and the other of X<sup>1</sup> and X<sup>2</sup> is CR<sup>9</sup>;

10 Z is NH, NHCO, NHSO<sub>2</sub>, NHCH<sub>2</sub>, CH<sub>2</sub>, CH<sub>2</sub>CH<sub>2</sub>, or CH=CH;

R<sup>1</sup>, R<sup>2</sup>, R<sup>3</sup>, R<sup>9</sup> and R<sup>10</sup> are independently H, alkyl, aryl, aralkyl, heterocycle, halogeno, NO<sub>2</sub>, CN, OH, alkoxy, aryloxy, (R''')<sub>n</sub>NH<sub>2</sub>, (R''')<sub>n</sub>NH-R', (R''')<sub>n</sub>N-(R')(R''), NH-aryl, N-(aryl)<sub>2</sub>, COOH, COO-R', COO-aryl, CONH<sub>2</sub>, CONH-R', CON-(R')(R''),  
15 CONH-aryl, CON-(aryl)<sub>2</sub>, SO<sub>3</sub>H, SO<sub>2</sub>NH<sub>2</sub>, CF<sub>3</sub>, CO-R', or CO-aryl, wherein alkyl, aryl, aralkyl and heterocycle groups may be further substituted with one or more groups selected from halogeno, NO<sub>2</sub>, CN, OH, O-methyl, NH<sub>2</sub>, COOH, CONH<sub>2</sub> and CF<sub>3</sub>;

20 R<sup>4</sup>, R<sup>5</sup>, R<sup>6</sup>, R<sup>7</sup>, and R<sup>8</sup> are independently from each other H, substituted or unsubstituted lower alkyl, halogeno, NO<sub>2</sub>, CN, OH, substituted or unsubstituted alkoxy, NH<sub>2</sub>, NH-R', N-(R')(R''), COOH, COO-R', CONH<sub>2</sub>, CONH-R', CON-(R')(R''), SO<sub>3</sub>H, SO<sub>2</sub>NH<sub>2</sub>, or CF<sub>3</sub>;

wherein R' R'' and R''' are each independently alkyl groups that may be the same or different and n is 0 or 1;

with the proviso that when R<sup>1</sup> and R<sup>2</sup> are H, X<sup>1</sup> is NH, X<sup>2</sup> is CH, and R<sup>3</sup> is H, the phenyl group is not

- 5           unsubstituted,  
            4-ethyl,  
            3-methyl,  
            3-(1,1,2,2- tetrafluoroethoxy),  
            3,4,5-trimethoxy,  
10       when the other groups R<sup>4</sup>-R<sup>8</sup> are H;  
          and pharmaceutically acceptable salts thereof.

*Description of the Preferred Embodiments*

- 15       As used herein the term "alkyl" includes both straight chain and branched alkyl groups having from 1 to 8 carbon atoms, e.g. methyl, ethyl propyl, isopropyl, butyl, isobutyl, tert-butyl, pentyl, hexyl etc and the term "lower alkyl" is similarly used for groups having from 1 to 4 carbon atoms.
- 20       The term "aryl" is used to include groups having from 6 to 10 carbon atoms, e.g. phenyl, naphthyl etc.

The term "aralkyl" is used as a conjunction of the terms alkyl and aryl as given above.

- 25       Preferred compounds of formula I are those bearing a mono-, di- or trisubstituted pyrrol radical, attached to the pyrimidine ring through one of the ring carbon atoms. Preferably, the pyrrol radical is a pyrrol-3-yl group (i.e. X<sup>1</sup> is CR<sup>9</sup> and X<sup>2</sup> is NR<sup>10</sup>, preferably NH) and is di- or tri-substituted.

The pyrrol group may be substituted by  $R^1$ ,  $R^2$ ,  $R^9$  and  $R^{10}$ . Preferably,  $R^1$ ,  $R^2$  and where appropriate  $R^9$  and  $R^{10}$  are independently H, alkyl, aryl, aralkyl, heterocycle, halogeno,  $NO_2$ , CN, OH, alkoxy, aryloxy,  $(R''')_nNH_2$ ,  $(R''')_nNH-R'$ ,  $(R''')_nN-(R')(R'')$ , NH-aryl, N-(aryl)<sub>2</sub>, COOH, COO- $R'$ , COO-aryl, CONH<sub>2</sub>, CONH- $R'$ , CON-  
5 (R')(R''), CONH-aryl, CON-(aryl)<sub>2</sub>, SO<sub>3</sub>H, SO<sub>2</sub>NH<sub>2</sub>, CF<sub>3</sub>, CO- $R'$ , or CO-aryl, wherein alkyl, aryl, aralkyl and heterocycle groups may be further substituted with one or more groups selected from halogeno,  $NO_2$ , CN, OH, O-methyl, NH<sub>2</sub>, COOH, CONH<sub>2</sub> and CF<sub>3</sub>. Most preferably  $R^{10}$  is H.

10 More preferably,  $R^1$ ,  $R^2$  and  $R^9$  are each independently selected from H, halogeno,  $NO_2$ , CN,  $(R''')_nN-(R')(R'')$ , CONH<sub>2</sub>, a C<sub>1-4</sub> alkyl group and a heterocyclic group. Preferably, at least one, more preferably at least two or three of  $R^1$ ,  $R^2$  and  $R^9$  are not hydrogen.

15 Preferably,  $R^1$  is H, CN, halogeno, nitro, alkylamino or a heterocyclic group. When  $R^1$  is halogeno, it is preferably selected from chloro or bromo. When  $R^1$  is alkylamino, it is preferably diethylaminomethyl or dimethylaminomethyl. When  $R^1$  is a heterocyclic group it is preferably morpholin-4-ylmethyl or 4-methyl-piperazin-1-ylmethyl. Most preferably,  $R^1$  is H or CN.

20

Even more preferably, when  $R^1$  is as preferably described,  $R^2$  and  $R^9$  are both lower alkyl, preferably methyl.

The group Z is preferably NH, NHSO<sub>2</sub> or NHCH<sub>2</sub>, most preferably NH.

25

The phenyl substituents  $R^4$ - $R^8$  are each independently selected from H, halogeno, nitro, amino, aminoalkyl, hydroxy, alkoxy, carbamoyl, sulfamyl, CN, N(R')(R''), C<sub>1-4</sub> alkyl and substituted C<sub>1-4</sub> alkyl.

More preferably,  $R^4$ ,  $R^5$ ,  $R^6$ ,  $R^7$ , and  $R^8$  are independently from each other H, unsubstituted lower alkyl, halogeno,  $\text{NO}_2$ , CN, OH,  $\text{N}-(R')(R'')$ , or  $\text{CF}_3$ ;

wherein  $R'$ ,  $R''$  and  $R'''$  are each independently alkyl groups that may be the same or different and n is 0 or 1;

5

Even more preferably,  $R^4$  to  $R^8$  are selected independently from H, F,  $\text{NH}_2$ ,  $\text{NO}_2$ , OH, Cl, Br, I, CN,  $\text{CH}_2\text{OH}$ ,  $\text{CF}_3$  and dimethylamino. Within the preferences for  $R^4$  to  $R^8$ ,  $R^4$  and  $R^8$  are most preferably hydrogen.

10 Thus, particularly preferred embodiments include 2-[N-(phenyl)]-4-(2,4-dimethylpyrrol-3-yl)pyrimidineamines in which the phenyl group is 2-, 3- or 4-substituted by at least one of H, F,  $\text{NH}_2$ ,  $\text{NO}_2$ , OH, Cl, Br, I, CN,  $\text{CH}_2\text{OH}$ ,  $\text{CF}_3$  or OMe.

15 Within this particular embodiment, the phenyl group is preferably mono-substituted by F,  $\text{NH}_2$ ,  $\text{NO}_2$ , OH, Cl, Br, I,  $\text{CH}_2\text{OH}$ , CN,  $\text{CF}_3$  or OMe at any of the 2,3 or 4-positions, or di-substituted by 2,4-difluoro, 3,5-difluoro, 3,4-difluoro, 2,4-dichloro, 3,5-dichloro, 3,4-dichloro or 4-chloro-3-trifluoromethyl.

20 Further particularly preferred embodiments include 2-[N-(phenyl)]-4-(3,5-dimethyl-1H-pyrrole-2-carbonitrile)pyrimidineamines in which the phenyl group is 2-, 3- or 4-substituted by at least one of F,  $\text{NH}(\text{CH}_3)_2$ ,  $\text{NO}_2$ , OH, Cl, Br, I or  $\text{CF}_3$ .

25 Within this particular embodiment, the phenyl group is preferably mono-substituted by F,  $\text{NH}(\text{CH}_3)_2$ ,  $\text{NO}_2$ , OH, I or  $\text{CF}_3$  at any of the 3 or 4-positions, or di-substituted by 4-methyl-3-nitro, 3-iodo-4-methyl, 4-chloro-3-methyl, 3-hydroxy-4-methyl, 4-fluoro-3-methyl or 4-methyl-3-fluoro.

Further more particularly preferable embodiments include;

- 2-[N-(phenyl)]-4-(2,4-Dimethyl-5-nitro-1H-pyrrol-3-yl)-pyrimidinamines wherein the phenyl group is preferably mono-substituted by F, NH(CH<sub>3</sub>)<sub>2</sub>, NO<sub>2</sub>, OH, I or CF<sub>3</sub> at the 4-position, preferably by a fluoro or NH(CH<sub>3</sub>)<sub>2</sub> group.
- 5 - 2-[N-(phenyl)]-4-(2,4-dimethyl-5-halogeno-1H-pyrrol-3-yl)-pyrimidinamines wherein the phenyl group is preferably mono-substituted by F, NH(CH<sub>3</sub>)<sub>2</sub>, NO<sub>2</sub>, OH, I or CF<sub>3</sub> at the 3 or 4-positions, preferably by a 4-fluoro or 3-nitro group, the halogeno group preferably being chloro or bromo.
- 10 - 2-[N-(phenyl)]-4-(2,4-dimethyl-5-dialkylaminoalkyl-1H-pyrrol-3-yl)-pyrimidinamines wherein the phenyl group is preferably mono-substituted by F, NH(CH<sub>3</sub>)<sub>2</sub>, NO<sub>2</sub>, OH, I or CF<sub>3</sub> at the 4-position, preferably by fluoro, the dialkylaminoalkyl group preferably being diethylaminomethyl or dimethylaminomethyl.
- 15 - 2-[N-(phenyl)]-4-(2,4-dimethyl-5-(heterocycle)-1H-pyrrol-3-yl)-pyrimidinamines wherein the phenyl group is preferably mono-substituted by F, NH(CH<sub>3</sub>)<sub>2</sub>, NO<sub>2</sub>, OH, I or CF<sub>3</sub> at the 4-position, preferably by fluoro, the heterocycle group preferably being 5-morpholin-4-ylmethyl or 4-methyl-piperazin-1-ylmethyl.
- 20 Most preferably, the compounds of the present invention are selected from;  
[4-(2,4-Dimethyl-1H-pyrrol-3-yl)-pyrimidin-2-yl]-(4-fluoro-phenyl)-amine  
[4-(2,4-Dimethyl-1H-pyrrol-3-yl)-pyrimidin-2-yl]-(3-nitro-phenyl)-amine  
[4-(2,4-Dimethyl-1H-pyrrol-3-yl)-pyrimidin-2-yl]-(4-iodo-phenyl)-amine  
25 (3,4-Difluoro-phenyl)-[4-(2,4-dimethyl-1H-pyrrol-3-yl)-pyrimidin-2-yl]-amine  
(4-Chloro-phenyl)-[4-(2,4-dimethyl-1H-pyrrol-3-yl)-pyrimidin-2-yl]-amine  
(3,5-Difluoro-phenyl)-[4-(2,4-dimethyl-1H-pyrrol-3-yl)-pyrimidin-2-yl]-amine  
4-[4-(2,4-Dimethyl-1H-pyrrol-3-yl)-pyrimidin-2-ylamino]-phenol  
3-[4-(2,4-Dimethyl-1H-pyrrol-3-yl)-pyrimidin-2-ylamino]-phenol

- (2,4-Difluoro-phenyl)-[4-(2,4-dimethyl-1H-pyrrol-3-yl)-pyrimidin-2-yl]-amine  
(2,4-Dichloro-phenyl)-[4-(2,4-dimethyl-1H-pyrrol-3-yl)-pyrimidin-2-yl]-amine  
(4-Chloro-3-trifluoromethyl-phenyl)-[4-(2,4-dimethyl-1H-pyrrol-3-yl)-pyrimidin-2-yl]-amine
- 5 [4-(2,4-Dimethyl-1H-pyrrol-3-yl)-pyrimidin-2-yl]-(4-trifluoromethyl-phenyl)-amine  
[4-(2,4-Dimethyl-1H-pyrrol-3-yl)-pyrimidin-2-yl]-(3-trifluoromethyl-phenyl)-amine  
(3-Chloro-phenyl)-[4-(2,4-dimethyl-1H-pyrrol-3-yl)-pyrimidin-2-yl]-amine  
N-[4-(2,4-Dimethyl-1H-pyrrol-3-yl)-pyrimidin-2-yl]-N',N'-dimethyl-benzene-1,4-diamine
- 10 (3-Chloro-4-iodo-phenyl)-[4-(2,4-dimethyl-1H-pyrrol-3-yl)-pyrimidin-2-yl]-amine  
[4-(2,4-Dimethyl-1H-pyrrol-3-yl)-pyrimidin-2-yl]-(3-fluoro-4-iodo-phenyl)-amine  
3,5-Dimethyl-4-[2-(3-nitro-phenylamino)-pyrimidin-4-yl]-1H-pyrrole-2-carbonitrile  
4-[2-(4-Fluoro-phenylamino)-pyrimidin-4-yl]-3,5-dimethyl-1H-pyrrole-2-carbonitrile  
4-[2-(4-Hydroxy-phenylamino)-pyrimidin-4-yl]-3,5-dimethyl-1H-pyrrole-2-
- 15 carbonitrile  
3,5-Dimethyl-4-[2-(4-trifluoromethyl-phenylamino)-pyrimidin-4-yl]-1H-pyrrole-2-carbonitrile  
4-[2-(4-Iodo-phenylamino)-pyrimidin-4-yl]-3,5-dimethyl-1H-pyrrole-2-carbonitrile  
4-[2-(3-Hydroxy-phenylamino)-pyrimidin-4-yl]-3,5-dimethyl-1H-pyrrole-2-
- 20 carbonitrile  
3,5-Dimethyl-4-[2-(4-methyl-3-nitro-phenylamino)-pyrimidin-4-yl]-1H-pyrrole-2-carbonitrile  
4-[2-(3-Iodo-4-methyl-phenylamino)-pyrimidin-4-yl]-3,5-dimethyl-1H-pyrrole-2-carbonitrile
- 25 4-[2-(4-Chloro-3-methyl-phenylamino)-pyrimidin-4-yl]-3,5-dimethyl-1H-pyrrole-2-carbonitrile  
4-[2-(3-Hydroxy-4-methyl-phenylamino)-pyrimidin-4-yl]-3,5-dimethyl-1H-pyrrole-2-carbonitrile

- 4-[2-(4-Fluoro-3-methyl-phenylamino)-pyrimidin-4-yl]-3,5-dimethyl-1H-pyrrole-2-carbonitrile
- 4-[2-(3-Fluoro-4-methyl-phenylamino)-pyrimidin-4-yl]-3,5-dimethyl-1H-pyrrole-2-carbonitrile
- 5 4-[2-(4-Dimethylamino-phenylamino)-pyrimidin-4-yl]-3,5-dimethyl-1H-pyrrole-2-carbonitrile
- 4-[2-(4-Fluoro-phenylamino)-pyrimidin-4-yl]-3,5-dimethyl-1H-pyrrole-2-carboxylic acid amide
- [4-(3,5-Dimethyl-1H-pyrrol-2-yl)-pyrimidin-2-yl]-(4-fluoro-phenyl)-amine
- 10 (4-Fluoro-phenyl)-[4-(1,2,4-trimethyl-1H-pyrrol-3-yl)-pyrimidin-2-yl]-amine
- [4-(2,4-Dimethyl-5-nitro-1H-pyrrol-3-yl)-pyrimidin-2-yl]-(4-fluoro-phenyl)-amine
- N-[4-(2,4-Dimethyl-5-nitro-1H-pyrrol-3-yl)-pyrimidin-2-yl]-N',N'-dimethyl-benzene-1,4-diamine
- [4-(5-Amino-2,4-dimethyl-1H-pyrrol-3-yl)-pyrimidin-2-yl]-(4-fluoro-phenyl)-amine
- 15 [4-(5-Bromo-2,4-dimethyl-1H-pyrrol-3-yl)-pyrimidin-2-yl]-(4-fluoro-phenyl)-amine
- [4-(5-Bromo-2,4-dimethyl-1H-pyrrol-3-yl)-pyrimidin-2-yl]-(3-nitro-phenyl)-amine
- [4-(5-Chloro-2,4-dimethyl-1H-pyrrol-3-yl)-pyrimidin-2-yl]-(4-fluoro-phenyl)-amine
- [4-(5-Diethylaminomethyl-2,4-dimethyl-1H-pyrrol-3-yl)-pyrimidin-2-yl]-(4-fluoro-phenyl)-amine
- 20 [4-(5-Dimethylaminomethyl-2,4-dimethyl-1H-pyrrol-3-yl)-pyrimidin-2-yl]-(4-fluoro-phenyl)-amine
- [4-(2,4-Dimethyl-5-morpholin-4-ylmethyl-1H-pyrrol-3-yl)-pyrimidin-2-yl]-(4-fluoro-phenyl)-amine
- {4-[2,4-Dimethyl-5-(4-methyl-piperazin-1-ylmethyl)-1H-pyrrol-3-yl]-pyrimidin-2-yl}-(4-fluoro-phenyl)-amine
- 25

The structures of the above-mentioned compounds are illustrated in Figure 1.

Particularly preferred compounds observed are those to be CDK inhibitors having  $IC_{50}$  for cdk2/cyclinE of less than  $5\mu M(\pm 0.05)$ , including;



- [4-(2,4-Dimethyl-1H-pyrrol-3-yl)-pyrimidin-2-yl]-(4-fluoro-phenyl)-amine  
[4-(2,4-Dimethyl-1H-pyrrol-3-yl)-pyrimidin-2-yl]-(3-nitro-phenyl)-amine  
[4-(2,4-Dimethyl-1H-pyrrol-3-yl)-pyrimidin-2-yl]-(4-iodo-phenyl)-amine  
(3,4-Difluoro-phenyl)-[4-(2,4-dimethyl-1H-pyrrol-3-yl)-pyrimidin-2-yl]-amine  
5 (4-Chloro-phenyl)-[4-(2,4-dimethyl-1H-pyrrol-3-yl)-pyrimidin-2-yl]-amine  
(3,5-Difluoro-phenyl)-[4-(2,4-dimethyl-1H-pyrrol-3-yl)-pyrimidin-2-yl]-amine  
4-[4-(2,4-Dimethyl-1H-pyrrol-3-yl)-pyrimidin-2-ylamino]-phenol  
3-[4-(2,4-Dimethyl-1H-pyrrol-3-yl)-pyrimidin-2-ylamino]-phenol  
[4-(2,4-Dimethyl-1H-pyrrol-3-yl)-pyrimidin-2-yl]-(4-trifluoromethyl-phenyl)-amine  
10 (3-Chloro-4-iodo-phenyl)-[4-(2,4-dimethyl-1H-pyrrol-3-yl)-pyrimidin-2-yl]-amine  
[4-(2,4-Dimethyl-1H-pyrrol-3-yl)-pyrimidin-2-yl]-(3-fluoro-4-iodo-phenyl)-amine  
3,5-Dimethyl-4-[2-(3-nitro-phenylamino)-pyrimidin-4-yl]-1H-pyrrole-2-carbonitrile  
4-[2-(4-Fluoro-phenylamino)-pyrimidin-4-yl]-3,5-dimethyl-1H-pyrrole-2-carbonitrile  
4-[2-(4-Hydroxy-phenylamino)-pyrimidin-4-yl]-3,5-dimethyl-1H-pyrrole-2-  
15 carbonitrile  
3,5-Dimethyl-4-[2-(4-trifluoromethyl-phenylamino)-pyrimidin-4-yl]-1H-pyrrole-2-  
carbonitrile  
4-[2-(4-Iodo-phenylamino)-pyrimidin-4-yl]-3,5-dimethyl-1H-pyrrole-2-carbonitrile  
4-[2-(3-Hydroxy-phenylamino)-pyrimidin-4-yl]-3,5-dimethyl-1H-pyrrole-2-  
20 carbonitrile  
3,5-Dimethyl-4-[2-(4-methyl-3-nitro-phenylamino)-pyrimidin-4-yl]-1H-pyrrole-2-  
carbonitrile  
4-[2-(3-Iodo-4-methyl-phenylamino)-pyrimidin-4-yl]-3,5-dimethyl-1H-pyrrole-2-  
carbonitrile  
25 4-[2-(4-Chloro-3-methyl-phenylamino)-pyrimidin-4-yl]-3,5-dimethyl-1H-pyrrole-2-  
carbonitrile  
4-[2-(3-Hydroxy-4-methyl-phenylamino)-pyrimidin-4-yl]-3,5-dimethyl-1H-pyrrole-2-  
carbonitrile

4-[2-(4-Fluoro-3-methyl-phenylamino)-pyrimidin-4-yl]-3,5-dimethyl-1H-pyrrole-2-carbonitrile

4-[2-(3-Fluoro-4-methyl-phenylamino)-pyrimidin-4-yl]-3,5-dimethyl-1H-pyrrole-2-carbonitrile

5 4-[2-(4-Dimethylamino-phenylamino)-pyrimidin-4-yl]-3,5-dimethyl-1H-pyrrole-2-carbonitrile

4-[2-(4-Fluoro-phenylamino)-pyrimidin-4-yl]-3,5-dimethyl-1H-pyrrole-2-carboxylic acid amide

(4-Fluoro-phenyl)-[4-(1,2,4-trimethyl-1H-pyrrol-3-yl)-pyrimidin-2-yl]-amine

10 [4-(2,4-Dimethyl-5-nitro-1H-pyrrol-3-yl)-pyrimidin-2-yl]-(4-fluoro-phenyl)-amine  
N-[4-(2,4-Dimethyl-5-nitro-1H-pyrrol-3-yl)-pyrimidin-2-yl]-N',N'-dimethyl-benzene-1,4-diamine

[4-(5-Bromo-2,4-dimethyl-1H-pyrrol-3-yl)-pyrimidin-2-yl]-(4-fluoro-phenyl)-amine

[4-(5-Bromo-2,4-dimethyl-1H-pyrrol-3-yl)-pyrimidin-2-yl]-(3-nitro-phenyl)-amine

15 [4-(5-Chloro-2,4-dimethyl-1H-pyrrol-3-yl)-pyrimidin-2-yl]-(4-fluoro-phenyl)-amine  
[4-(5-Diethylaminomethyl-2,4-dimethyl-1H-pyrrol-3-yl)-pyrimidin-2-yl]-(4-fluoro-phenyl)-amine

[4-(5-Dimethylaminomethyl-2,4-dimethyl-1H-pyrrol-3-yl)-pyrimidin-2-yl]-(4-fluoro-phenyl)-amine

20 [4-(2,4-Dimethyl-5-morpholin-4-ylmethyl-1H-pyrrol-3-yl)-pyrimidin-2-yl]-(4-fluoro-phenyl)-amine, and

{4-[2,4-Dimethyl-5-(4-methyl-piperazin-1-ylmethyl)-1H-pyrrol-3-yl]-pyrimidin-2-yl}-(4-fluoro-phenyl)-amine.

25 Of these compounds, more preferred are those to be CDK inhibitors having IC<sub>50</sub> for cdk2/cyclinE of less than 1 μM(±0.05), including;

[4-(2,4-Dimethyl-1H-pyrrol-3-yl)-pyrimidin-2-yl]-(4-fluoro-phenyl)-amine

[4-(2,4-Dimethyl-1H-pyrrol-3-yl)-pyrimidin-2-yl]-(3-nitro-phenyl)-amine

- [4-(2,4-Dimethyl-1H-pyrrol-3-yl)-pyrimidin-2-yl]-(4-iodo-phenyl)-amine  
[4-(2,4-Dimethyl-1H-pyrrol-3-yl)-pyrimidin-2-yl]-(4-trifluoromethyl-phenyl)-amine  
3,5-Dimethyl-4-[2-(3-nitro-phenylamino)-pyrimidin-4-yl]-1H-pyrrole-2-carbonitrile  
4-[2-(4-Fluoro-phenylamino)-pyrimidin-4-yl]-3,5-dimethyl-1H-pyrrole-2-carbonitrile  
5 4-[2-(4-Hydroxy-phenylamino)-pyrimidin-4-yl]-3,5-dimethyl-1H-pyrrole-2-carbonitrile  
3,5-Dimethyl-4-[2-(4-trifluoromethyl-phenylamino)-pyrimidin-4-yl]-1H-pyrrole-2-carbonitrile  
4-[2-(4-Iodo-phenylamino)-pyrimidin-4-yl]-3,5-dimethyl-1H-pyrrole-2-carbonitrile  
10 4-[2-(3-Hydroxy-phenylamino)-pyrimidin-4-yl]-3,5-dimethyl-1H-pyrrole-2-carbonitrile  
3,5-Dimethyl-4-[2-(4-methyl-3-nitro-phenylamino)-pyrimidin-4-yl]-1H-pyrrole-2-carbonitrile  
4-[2-(3-Iodo-4-methyl-phenylamino)-pyrimidin-4-yl]-3,5-dimethyl-1H-pyrrole-2-carbonitrile  
15 4-[2-(4-Chloro-3-methyl-phenylamino)-pyrimidin-4-yl]-3,5-dimethyl-1H-pyrrole-2-carbonitrile  
4-[2-(3-Hydroxy-4-methyl-phenylamino)-pyrimidin-4-yl]-3,5-dimethyl-1H-pyrrole-2-carbonitrile  
20 4-[2-(4-Fluoro-3-methyl-phenylamino)-pyrimidin-4-yl]-3,5-dimethyl-1H-pyrrole-2-carbonitrile  
4-[2-(3-Fluoro-4-methyl-phenylamino)-pyrimidin-4-yl]-3,5-dimethyl-1H-pyrrole-2-carbonitrile  
4-[2-(4-Dimethylamino-phenylamino)-pyrimidin-4-yl]-3,5-dimethyl-1H-pyrrole-2-carbonitrile  
25 4-[2-(4-Fluoro-phenylamino)-pyrimidin-4-yl]-3,5-dimethyl-1H-pyrrole-2-carboxylic acid amide  
[4-(2,4-Dimethyl-5-nitro-1H-pyrrol-3-yl)-pyrimidin-2-yl]-(4-fluoro-phenyl)-amine

- N-[4-(2,4-Dimethyl-5-nitro-1H-pyrrol-3-yl)-pyrimidin-2-yl]-N',N'-dimethyl-benzene-1,4-diamine
- [4-(5-Bromo-2,4-dimethyl-1H-pyrrol-3-yl)-pyrimidin-2-yl]-(4-fluoro-phenyl)-amine
- [4-(5-Bromo-2,4-dimethyl-1H-pyrrol-3-yl)-pyrimidin-2-yl]-(3-nitro-phenyl)-amine
- 5 [4-(5-Chloro-2,4-dimethyl-1H-pyrrol-3-yl)-pyrimidin-2-yl]-(4-fluoro-phenyl)-amine
- [4-(5-Diethylaminomethyl-2,4-dimethyl-1H-pyrrol-3-yl)-pyrimidin-2-yl]-(4-fluoro-phenyl)-amine
- [4-(5-Dimethylaminomethyl-2,4-dimethyl-1H-pyrrol-3-yl)-pyrimidin-2-yl]-(4-fluoro-phenyl)-amine, and
- 10 [4-(2,4-Dimethyl-5-morpholin-4-ylmethyl-1H-pyrrol-3-yl)-pyrimidin-2-yl]-(4-fluoro-phenyl)-amine.

Of these, even more preferred are compounds are those having  $IC_{50}$  for cdk2/cyclinE of less than  $0.5\mu M$  ( $\pm 0.05$ ), being;

- 15 [4-(2,4-Dimethyl-1H-pyrrol-3-yl)-pyrimidin-2-yl]-(3-nitro-phenyl)-amine
- [4-(2,4-Dimethyl-1H-pyrrol-3-yl)-pyrimidin-2-yl]-(4-iodo-phenyl)-amine
- 3,5-Dimethyl-4-[2-(3-nitro-phenylamino)-pyrimidin-4-yl]-1H-pyrrole-2-carbonitrile
- 4-[2-(4-Fluoro-phenylamino)-pyrimidin-4-yl]-3,5-dimethyl-1H-pyrrole-2-carbonitrile
- 20 4-[2-(4-Hydroxy-phenylamino)-pyrimidin-4-yl]-3,5-dimethyl-1H-pyrrole-2-carbonitrile
- 3,5-Dimethyl-4-[2-(4-trifluoromethyl-phenylamino)-pyrimidin-4-yl]-1H-pyrrole-2-carbonitrile
- 4-[2-(4-Iodo-phenylamino)-pyrimidin-4-yl]-3,5-dimethyl-1H-pyrrole-2-carbonitrile
- 25 4-[2-(3-Hydroxy-phenylamino)-pyrimidin-4-yl]-3,5-dimethyl-1H-pyrrole-2-carbonitrile
- 3,5-Dimethyl-4-[2-(4-methyl-3-nitro-phenylamino)-pyrimidin-4-yl]-1H-pyrrole-2-carbonitrile

4-[2-(3-Hydroxy-4-methyl-phenylamino)-pyrimidin-4-yl]-3,5-dimethyl-1H-pyrrole-2-carbonitrile

4-[2-(4-Fluoro-3-methyl-phenylamino)-pyrimidin-4-yl]-3,5-dimethyl-1H-pyrrole-2-carbonitrile

- 5 4-[2-(4-Fluoro-phenylamino)-pyrimidin-4-yl]-3,5-dimethyl-1H-pyrrole-2-carboxylic acid amide

[4-(2,4-Dimethyl-5-nitro-1H-pyrrol-3-yl)-pyrimidin-2-yl]-(4-fluoro-phenyl)-amine

[4-(5-Bromo-2,4-dimethyl-1H-pyrrol-3-yl)-pyrimidin-2-yl]-(3-nitro-phenyl)-amine,  
and

- 10 [4-(5-Dimethylaminomethyl-2,4-dimethyl-1H-pyrrol-3-yl)-pyrimidin-2-yl]-(4-fluoro-phenyl)-amine.

The following compounds are observed to be particularly effective anti-proliferative agents, as demonstrated by cell-based assays:

- 15 [4-(2,4-Dimethyl-1H-pyrrol-3-yl)-pyrimidin-2-yl]-(3-nitro-phenyl)-amine  
[4-(2,4-Dimethyl-1H-pyrrol-3-yl)-pyrimidin-2-yl]-(4-trifluoromethyl-phenyl)-amine  
(3-Chloro-4-iodo-phenyl)-[4-(2,4-dimethyl-1H-pyrrol-3-yl)-pyrimidin-2-yl]-amine  
3,5-Dimethyl-4-[2-(3-nitro-phenylamino)-pyrimidin-4-yl]-1H-pyrrole-2-carbonitrile  
4-[2-(4-Fluoro-phenylamino)-pyrimidin-4-yl]-3,5-dimethyl-1H-pyrrole-2-carbonitrile  
20 4-[2-(4-Hydroxy-phenylamino)-pyrimidin-4-yl]-3,5-dimethyl-1H-pyrrole-2-carbonitrile  
3,5-Dimethyl-4-[2-(4-trifluoromethyl-phenylamino)-pyrimidin-4-yl]-1H-pyrrole-2-carbonitrile  
4-[2-(4-Iodo-phenylamino)-pyrimidin-4-yl]-3,5-dimethyl-1H-pyrrole-2-carbonitrile  
25 4-[2-(3-Hydroxy-phenylamino)-pyrimidin-4-yl]-3,5-dimethyl-1H-pyrrole-2-carbonitrile  
3,5-Dimethyl-4-[2-(4-methyl-3-nitro-phenylamino)-pyrimidin-4-yl]-1H-pyrrole-2-carbonitrile

- 4-[2-(3-Iodo-4-methyl-phenylamino)-pyrimidin-4-yl]-3,5-dimethyl-1H-pyrrole-2-carbonitrile
- 4-[2-(4-Chloro-3-methyl-phenylamino)-pyrimidin-4-yl]-3,5-dimethyl-1H-pyrrole-2-carbonitrile
- 5 4-[2-(3-Hydroxy-4-methyl-phenylamino)-pyrimidin-4-yl]-3,5-dimethyl-1H-pyrrole-2-carbonitrile
- 4-[2-(4-Fluoro-3-methyl-phenylamino)-pyrimidin-4-yl]-3,5-dimethyl-1H-pyrrole-2-carbonitrile
- 4-[2-(3-Fluoro-4-methyl-phenylamino)-pyrimidin-4-yl]-3,5-dimethyl-1H-pyrrole-2-carbonitrile
- 10 4-[2-(4-Dimethylamino-phenylamino)-pyrimidin-4-yl]-3,5-dimethyl-1H-pyrrole-2-carbonitrile
- 4-[2-(4-Fluoro-phenylamino)-pyrimidin-4-yl]-3,5-dimethyl-1H-pyrrole-2-carboxylic acid amide
- 15 (4-Fluoro-phenyl)-[4-(1,2,4-trimethyl-1H-pyrrol-3-yl)-pyrimidin-2-yl]-amine
- [4-(2,4-Dimethyl-5-nitro-1H-pyrrol-3-yl)-pyrimidin-2-yl]-(4-fluoro-phenyl)-amine
- N-[4-(2,4-Dimethyl-5-nitro-1H-pyrrol-3-yl)-pyrimidin-2-yl]-N',N'-dimethyl-benzene-1,4-diamine
- [4-(5-Bromo-2,4-dimethyl-1H-pyrrol-3-yl)-pyrimidin-2-yl]-(3-nitro-phenyl)-amine
- 20 [4-(5-Diethylaminomethyl-2,4-dimethyl-1H-pyrrol-3-yl)-pyrimidin-2-yl]-(4-fluoro-phenyl)-amine
- [4-(5-Dimethylaminomethyl-2,4-dimethyl-1H-pyrrol-3-yl)-pyrimidin-2-yl]-(4-fluoro-phenyl)-amine, and
- {4-[2,4-Dimethyl-5-(4-methyl-piperazin-1-ylmethyl)-1H-pyrrol-3-yl]-pyrimidin-2-yl}-(4-fluoro-phenyl)-amine.
- 25

The compounds of formula I have been found to possess anti-proliferative activity and are therefore believed to be of use in the treatment of proliferative disorders such as cancers, leukaemias and other disorders associated with uncontrolled cellular

proliferation such as psoriasis and restenosis. As defined herein, an anti-proliferative effect within the scope of the present invention may be demonstrated by the ability to inhibit cell proliferation in an *in vitro* whole cell assay, for example using any of the cell lines A549, HT29, Saos-2, HeLa or MCF-7, or by showing inhibition of a CDK  
5 enzyme (such as CDK2 or CDK4) in an appropriate assay. These assays, including methods for their performance, are described in more detail in Example 3. Using such cell line and enzymes assays it may be determined whether a compound is anti-proliferative in the context of the present invention.

10 Without wishing to be bound by theory, the compounds of the present invention are believed to exert their anti-proliferative effect in a non-protein kinase C (PKC) dependent manner. Many of the compounds inhibit cyclin-dependent kinase enzymes (CDKs) that have been shown to be involved in cell cycle control. These CDKs include CDK2 and CDK4 and particularly their respective interactions with cyclin E  
15 and cyclin D1. These compounds of the present invention are further believed to be advantageous in being selective for CDK enzymes implicated in proliferative diseases. By the term "selective" it is meant that although possible having some inhibitory effect on another enzyme (such as PKC), the compound is preferentially effective against an enzyme implicated in proliferative diseases.

20 The compounds of the invention may inhibit any of the steps or stages in the cell cycle, for example, formation of the nuclear envelope, exit from the quiescent phase of the cell cycle (G0), G1 progression, chromosome decondensation, nuclear envelope breakdown, START, initiation of DNA replication, progression of DNA replication,  
25 termination of DNA replication, centrosome duplication, G2 progression, activation of mitotic or meiotic functions, chromosome condensation, centrosome separation, microtubule nucleation, spindle formation and function, interactions with microtubule motor proteins, chromatid separation and segregation, inactivation of mitotic functions, formation of contractile ring, and cytokinesis functions. In particular, the

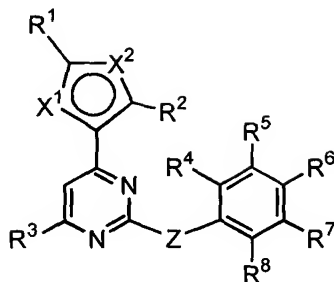
compounds of the invention may influence certain gene functions such as chromatin binding, formation of replication complexes, replication licensing, phosphorylation or other secondary modification activity, proteolytic degradation, microtubule binding, actin binding, septin binding, microtubule organising centre nucleation activity and  
5 binding to components of cell cycle signalling pathways.

A further embodiment of the present invention therefore relates to the use of one or more compounds of formula I in the treatment of proliferative disorders. Preferably, the proliferative disorder is a cancer or leukaemia. The term proliferative disorder is  
10 used herein in a broad sense to include any disorder that requires control of the cell cycle, for example cardiovascular disorders such as restenosis and cardiomyopathy, auto-immune disorders such as glomerulonephritis and rheumatoid arthritis, dermatological disorders such as psoriasis, anti-inflammatory, anti-fungal, antiparasitic disorders such as malaria, emphysema and alopecia. In these disorders,  
15 the compounds of the present invention may induce apoptosis or maintain stasis within the desired cells as required.

In a particularly preferred embodiment, the invention relates to the use of one or more compounds of formula I in the treatment of a CDK dependent or sensitive disorder.  
20 CDK dependent disorders are associated with an above normal level of activity of one or more CDK enzymes. Such disorders preferably associated with an abnormal level of activity of CDK2 and/or CDK4. A CDK sensitive disorder is a disorder in which an aberration in the CDK level is not the primary cause, but is downstream of the primary metabolic aberration. In such scenarios, CDK2 and/or CDK4 can be said to  
25 be part of the sensitive metabolic pathway and CDK inhibitors may therefore be active in treating such disorders. Such disorders are preferably cancer or leukaemic disorders.

A second aspect of the present invention relates to the use of a compound of formula





II

5 wherein:

one of X<sup>1</sup> and X<sup>2</sup> is NH and the other of X<sup>1</sup> and X<sup>2</sup> is CR<sup>9</sup>;

Z is NH, NHCO, NHSO<sub>2</sub>, NHCH<sub>2</sub>, CH<sub>2</sub>, CH<sub>2</sub>CH<sub>2</sub>, or CH=CH;

- 10 R<sup>1</sup>, R<sup>2</sup>, R<sup>3</sup> and R<sup>9</sup> are independently H, alkyl, aryl, aralkyl, heterocycle, halogeno, NO<sub>2</sub>, CN, OH, alkoxy, aryloxy, NH<sub>2</sub>, NH-R', N-(R')(R''), NH-aryl, N-(aryl)<sub>2</sub>, COOH, COO-R', COO-aryl, CONH<sub>2</sub>, CONH-R', CON-(R')(R''), CONH-aryl, CON-(aryl)<sub>2</sub>, SO<sub>3</sub>H, SO<sub>2</sub>NH<sub>2</sub>, CF<sub>3</sub>, CO-R', or CO-aryl, wherein alkyl, aryl, aralkyl and heterocycle groups may be further substituted with one or more groups selected from halogeno,  
15 NO<sub>2</sub>, CN, OH, O-methyl, NH<sub>2</sub>, COOH, CONH<sub>2</sub> and CF<sub>3</sub>;

- R<sup>4</sup>, R<sup>5</sup>, R<sup>6</sup>, R<sup>7</sup>, and R<sup>8</sup> are independently from each other H, substituted or unsubstituted lower alkyl, halogeno, NO<sub>2</sub>, CN, OH, substituted or unsubstituted alkoxy, NH<sub>2</sub>, NH-R', N-(R')(R''), COOH, COO-R', CONH<sub>2</sub>, CONH-R', CON-  
20 (R')(R''), SO<sub>3</sub>H, SO<sub>2</sub>NH<sub>2</sub>, or CF<sub>3</sub>;

wherein R' and R'' are each independently alkyl groups that may be the same or different;

with the proviso that when  $R^1$  and  $R^2$  are H,  $X^1$  is NH,  $X^2$  is CH, and  $R^3$  is H, the phenyl group is not

3-(1,1,2,2- tetrafluoroethoxy), or

3,4,5-trimethoxy,

- 5 when the other groups  $R^4$ - $R^8$  are H;  
and pharmaceutically acceptable salts thereof;  
in the manufacture of a medicament for use in the treatment of a proliferative disease.

10 The term “proliferative disorder” has been previously discussed and the same definition applies to the second aspect of the invention.

A further aspect of the present invention relates to the use of the compounds of formula II in the manufacture of a medicament for use in the treatment of antiviral infections. Such viral infections include VZV, HSV type 1 and 2 and HIV.  
15 Preferably, the compounds are of use in the treatment of HIV and HIV related disorders.

The preferred embodiments of these further aspects of the invention are identical to those described above in respect of the first aspect.

20 In a particularly preferred embodiment, the one or more compounds of the invention are administered in combination with one or more other anticancer agents. In such cases, the compounds of the invention may be administered consecutively, simultaneously or sequentially with the one or more other anticancer agents.

25 As used herein the phrase “manufacture of a medicament” includes the use of a compound of formula I directly as the medicament in addition to its use in a screening programme for further anti-proliferative agents or in any stage of the manufacture of such a medicament.

The compounds of the present invention (first and seconds aspects) can be present as salts or esters, in particular pharmaceutically acceptable salts or esters.

5 Pharmaceutically acceptable salts of the compounds of the invention (first and seconds aspects) include suitable acid addition or base salts thereof. A review of suitable pharmaceutical salts may be found in Berge et al, J Pharm Sci, 66, 1-19 (1977). Salts are formed, for example with strong inorganic acids such as mineral acids, e.g. sulphuric acid, phosphoric acid or hydrohalic acids; with strong organic  
10 carboxylic acids, such as alkanecarboxylic acids of 1 to 4 carbon atoms which are unsubstituted or substituted (e.g., by halogen), such as acetic acid; with saturated or unsaturated dicarboxylic acids, for example oxalic, malonic, succinic, maleic, fumaric, phthalic or tetraphthalic; with hydroxycarboxylic acids, for example ascorbic, glycolic, lactic, malic, tartaric or citric acid; with aminoacids, for example aspartic or  
15 glutamic acid; with benzoic acid; or with organic sulfonic acids, such as (C<sub>1</sub>-C<sub>4</sub>)-alkyl- or aryl-sulfonic acids which are unsubstituted or substituted (for example, by a halogen) such as methane- or p-toluene sulfonic acid.

Esters are formed either using organic acids or alcohols/hydroxides, depending on the  
20 functional group being esterified. Organic acids include carboxylic acids, such as alkanecarboxylic acids of 1 to 12 carbon atoms which are unsubstituted or substituted (e.g., by halogen), such as acetic acid; with saturated or unsaturated dicarboxylic acid, for example oxalic, malonic, succinic, maleic, fumaric, phthalic or tetraphthalic; with hydroxycarboxylic acids, for example ascorbic, glycolic, lactic, malic, tartaric or citric  
25 acid; with aminoacids, for example aspartic or glutamic acid; with benzoic acid; or with organic sulfonic acids, such as (C<sub>1</sub>-C<sub>4</sub>)-alkyl- or aryl-sulfonic acids which are unsubstituted or substituted (for example, by a halogen) such as methane- or p-toluene sulfonic acid. Suitable hydroxides include inorganic hydroxides, such as sodium hydroxide, potassium hydroxide, calcium hydroxide, aluminium hydroxide. Alcohols

include alkanealcohols of 1-12 carbon atoms which may be unsubstituted or substituted, e.g. by a halogen).

5 In all aspects of the present invention previously discussed, the invention includes, where appropriate all enantiomers and tautomers of compounds of formula I. The man skilled in the art will recognise compounds that possess an optical properties (one or more chiral carbon atoms) or tautomeric characteristics. The corresponding enantiomers and/or tautomers may be isolated/prepared by methods known in the art.

10 The invention furthermore relates to the compounds of or of use in the present invention (first and seconds aspects) in their various crystalline forms, polymorphic forms and (an)hydrous forms. It is well established within the pharmaceutical industry that chemical compounds may be isolated in any of such forms by slightly varying the method of purification and or isolation form the solvents used in the  
15 synthetic preparation of such compounds.

The invention further includes the compounds (first and seconds aspects) of or of use in the present invention in prodrug form. Such prodrugs are generally compounds of formula I wherein one or more appropriate groups have been modified such that the  
20 modification may be reversed upon administration to a human or mammalian subject. Such reversion is usually performed by an enzyme naturally present in such subject, though it is possible for a second agent to be administered together with such a prodrug in order to perform the reversion in vivo. Examples of such modifications include ester (for example, any of those described above), wherein the reversion may  
25 be carried out be an esterase etc. Other such systems will be well known to those skilled in the art.

The present invention also encompasses pharmaceutical compositions comprising the compounds of the invention (first and seconds aspects). In this regard, and in

particular for human therapy, even though the compounds of the present invention (including their pharmaceutically acceptable salts, esters and pharmaceutically acceptable solvates) can be administered alone, they will generally be administered in admixture with a pharmaceutical carrier, excipient or diluent selected with regard to the intended route of administration and standard pharmaceutical practice.

Thus, the present invention also relates to pharmaceutical compositions comprising one or more compounds of formula I or II or pharmaceutically acceptable salts or esters thereof, together with at least one pharmaceutically acceptable excipient, diluent or carrier.

By way of example, in the pharmaceutical compositions of the present invention, the compounds of the invention may be admixed with any suitable binder(s), lubricant(s), suspending agent(s), coating agent(s), and/or solubilising agent(s). Examples of such suitable excipients for the various different forms of pharmaceutical compositions described herein may be found in the "Handbook of Pharmaceutical Excipients, 2<sup>nd</sup> Edition, (1994), Edited by A Wade and PJ Weller.

The pharmaceutical compositions of the present invention may be adapted for oral, rectal, vaginal, parenteral, intramuscular, intraperitoneal, intraarterial, intrathecal, intrabronchial, subcutaneous, intradermal, intravenous, nasal, buccal or sublingual routes of administration.

For oral administration, particular use is made of compressed tablets, pills, tablets, gellules, drops, and capsules. Preferably, these compositions contain from 1 to 250 mg and more preferably from 10-100 mg, of active ingredient per dose.

Other forms of administration comprise solutions or emulsions which may be injected intravenously, intraarterially, intrathecally, subcutaneously, intradermally,

intraperitoneally or intramuscularly, and which are prepared from sterile or sterilisable solutions. The pharmaceutical compositions of the present invention may also be in form of suppositories, pessaries, suspensions, emulsions, lotions, ointments, creams, gels, sprays, solutions or dusting powders.

5

An alternative means of transdermal administration is by use of a skin patch. For example, the active ingredient can be incorporated into a cream consisting of an aqueous emulsion of polyethylene glycols or liquid paraffin. The active ingredient can also be incorporated, at a concentration of between 1 and 10% by weight, into an ointment consisting of a white wax or white soft paraffin base together with such stabilisers and preservatives as may be required.

10

Injectable forms may contain between 10 - 1000 mg, preferably between 10 - 250 mg, of active ingredient per dose.

15

Compositions may be formulated in unit dosage form, i.e., in the form of discrete portions containing a unit dose, or a multiple or sub-unit of a unit dose.

20

A person of ordinary skill in the art can easily determine an appropriate dose of one of the instant compositions to administer to a subject without undue experimentation. Typically, a physician will determine the actual dosage which will be most suitable for an individual patient and it will vary with the age, weight and response of the particular patient. The dosages disclosed herein are exemplary of the average case. There can of course be individual instances where higher or lower dosage ranges are merited, and such are within the scope of this invention.

25

In an exemplary embodiment, one or more doses of 10 to 150 mg/day will be administered to the patient for the treatment of malignancy.

The pharmaceutical compositions of the invention may further comprise one or more additional anticancer agents, for example, existing anticancer drugs available on the market.

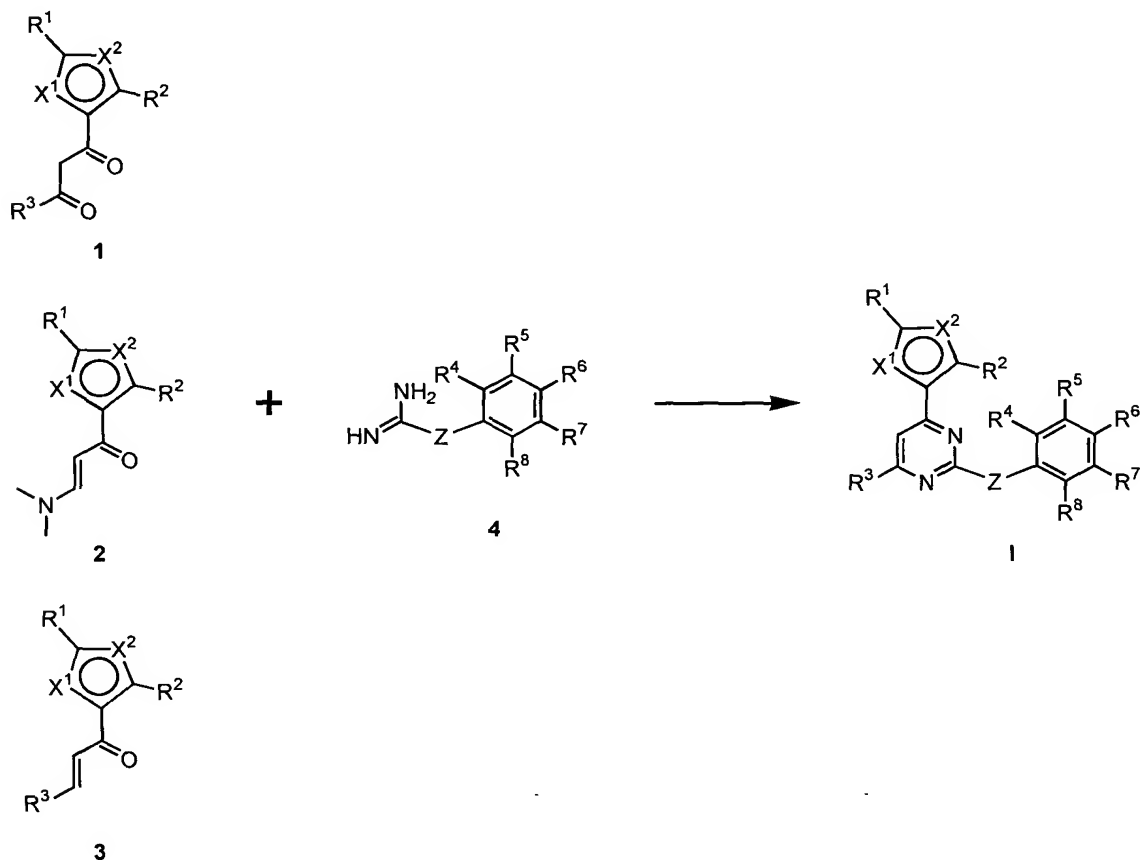
5 Anticancer drugs in general are more effective when used in combination. In particular, combination therapy is desirable in order to avoid an overlap of major toxicities, mechanism of action and resistance mechanism(s). Furthermore, it is also desirable to administer most drugs at their maximum tolerated doses with minimum time intervals between such doses. The major advantages of combining  
10 chemotherapeutic drugs are that it may promote additive or possible synergistic effects through biochemical interactions and also may decrease the emergence of resistance in early tumor cells which would have been otherwise responsive to initial chemotherapy with a single agent. An example of the use of biochemical interactions in selecting drug combinations is demonstrated by the administration of leucovorin to increase the  
15 binding of an active intracellular metabolite of 5-fluorouracil to its target, thymidylate synthase, thus increasing its cytotoxic effects.

Numerous combinations are used in current treatments of cancer and leukemia. A more extensive review of medical practices may be found in "Oncologic Therapies"  
20 edited by E. E. Vokes and H. M. Golomb, published by Springer.

Beneficial combinations may be suggested by studying the growth inhibitory activity of the test compounds with agents known or suspected of being valuable in the treatment of a particular cancer initially or cell lines derived from that cancer. This  
25 procedure can also be used to determine the order of administration of the agents, i.e. before, simultaneously, or after delivery. Such scheduling may be a feature of all the cycle acting agents identified herein.

The compounds of this invention (I) can be synthesised, for example, by an adaptation of the Traube synthesis (A.R. Katritzky, I. Taher, *Can. J. Chem.* 1986, 64, 2087 and references cited therein), *i.e.* by condensation between 1,3-dicarbonyl compounds **1** or acrylates **2** or **3**, and amidine **4**, as shown in *Scheme 1*.

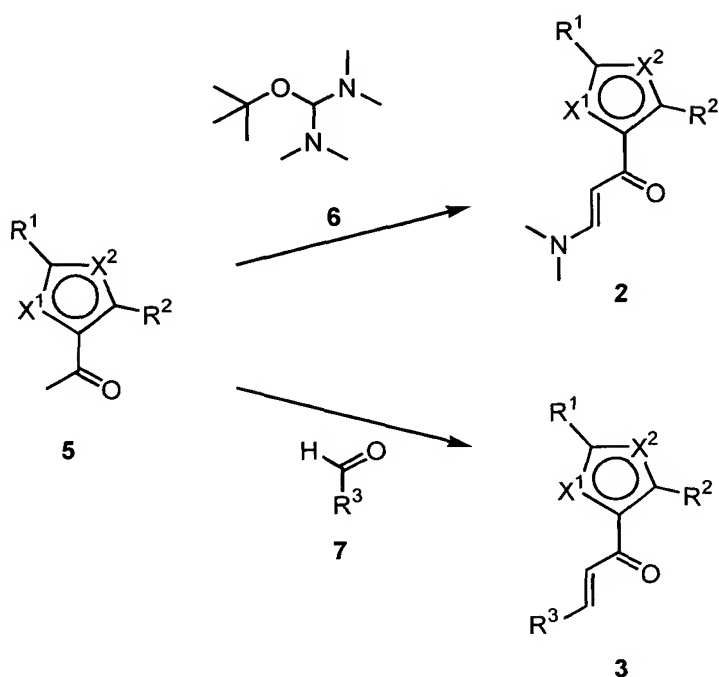
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*Scheme 1*

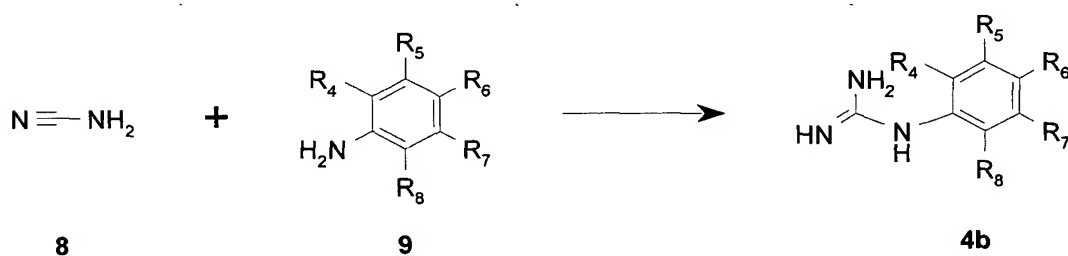
The dicarbonyl compounds **I** in turn can be prepared by many methods known in the art (J. March, *In: Advanced Organic Chemistry: Reactions, Mechanism, and Structure*, 4<sup>th</sup> Ed., John Wiley & Sons, Inc., New York, 1992, p. 1283). Acrylates **2** and **3**, which are particularly suitable for the purposes of this invention, are obtained from heterocyclic methyl ketones **5** by condensation with *tert*-butoxybis(dimethylamino)methane **6** (*Scheme 2*).





Scheme 2

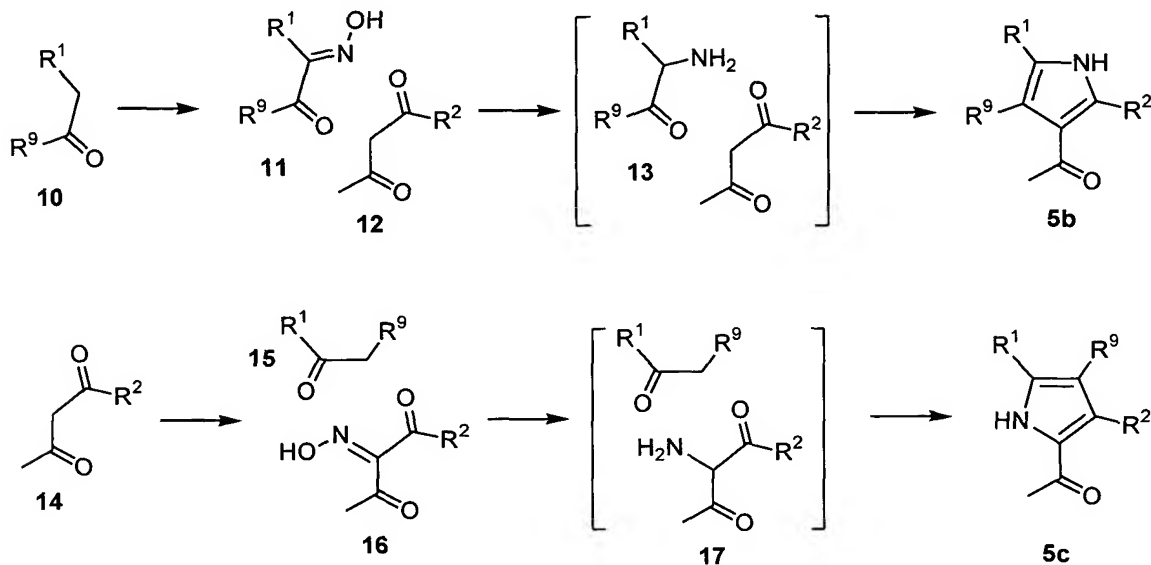
The diamino compounds **4** will be amidines **4a** or guanidines **4b**, depending on the definition of Z in general structure **I**. Amidines (HN=CRNH<sub>2</sub>) can be obtained from readily available amine precursors by condensation with *e.g.* ketenimines, or by addition of ammonia to suitable nitriles or imidates. Guanidines **4b** (Scheme 3) can be elaborated by a number of methods known in the art. For the purposes of this invention, the most useful route is amination of cyanamide **8** with anilines **9**.



Scheme 3

In the case where **5** is a pyrrole, two systems can apply (refer Scheme 4), *i.e.* the acetyl group which is used to generate the pyrimidine precursors **2** and **3** is either in the

pyrrole 3-position (**5**:  $X^1 = CR^9$ ,  $X^2 = NH$ ; *i.e.* structure **5b**) or in the pyrrole 2-position ( $X^1 = NH$ ,  $X^2 = CR^9$ ; *i.e.* structure **5c**).



5 Scheme 4

In both cases the pyrrole rings can be assembled using methods known in the art. Particularly relevant is a modification of the Knorr synthesis (refer, *e.g.* J. A. Joule, G. F. Smith, *Heterocyclic Chemistry*, 2<sup>nd</sup> Ed., Van Nostrand Reinhold (UK) Co. Ltd., 1978, pp.213-215). For the pyrrol-3-yl system, activated (*i.e.*  $R^1 = COOEt$ , CN, *etc.*) carbonyl compounds **10** are first nitrosylated. The resulting oximes **11** are condensed with dicarbonyl compounds **12** in the presence of *e.g.* zinc-acetic acid or aqueous dithionite, with formation of the reactive  $\alpha$ -aminocarbonyl intermediate **13**. The  $R^1$  substituent (*e.g.*  $COOEt$ , CN) in the resulting 3-acetylpyrroles **5b** can be further manipulated, either directly, or in the context of intermediates **2** or **3**, or in the pyrrolopyrimidine products **I**. Thus decarboxylation ( $R^1 = COOEt$ ) will give products with  $R^1 = H$ , oxidation ( $R^1 = CN$ ) will afford products with  $R^1 = CONH_2$ , *etc.* Furthermore, products with  $R^1 = H$  can be transformed into various derivatives, particularly through electrophilic substitution. Thus derivatives where  $R^1$  is, for example, a halogen, nitro, amino, alkyl, alkylamino, *etc.*, group can be obtained

readily. In the case of the pyrrol-2-yl system an analogous situation arises, here an activating group needs to be present in the carbonyl component 15 (*e.g.*  $R^9 = \text{COOEt}$ , CN, *etc.*). This is condensed with oximes 16 (derived from dicarbonyl compounds 14), again with formation of the intermediate 17. The  $R^9$  substituent in products 5c or derivatives can be manipulated in the same way as the  $R^1$  group in the pyrrol-3-yl system discussed above.

Alternatively, compounds of general structure I can be obtained from suitable pyrimidine precursors directly, *e.g.* from 2,4-disubstituted (halogen, amine, *etc.*) pyrimidines by successive substitution reactions.

The present invention will now be described by way of example and with reference to the following figure:

Figure 1 shows the chemical structure of compounds according to the invention.

### Examples

#### Abbreviations

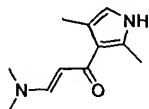
LC-MS, liquid chromatography-mass spectrometry; NMR, nuclear magnetic resonance spectroscopy; r.t. room temperature; PE, petroleum ether (40-60 °C boiling fraction); DMSO, dimethylsulfoxide.

#### General

NMR spectra were recorded using a Bruker DPX-300 instrument. Chemical shifts are reported in ppm ( $\delta$ ) from tetramethylsilane. EM Kieselgel 60 (0.040-0.063 mm) was used for flash column chromatography. Melting points were determined with a LEICA testo-720 electrothermometer and are uncorrected. Compound numbers are shown in brackets, where appropriate.

**Example 1**

*3-Dimethylamino-1-(2,4-dimethyl-1H-pyrrol-3-yl)-propenone*



A mixture of 1-(2,4-dimethyl-1H-pyrrol-3-yl)-ethanone (2 g, 15 mmol) in 5 mL of  
5 1,1-bis-dimethylamino-3,3-dimethyl-butan-2-one was heated at 100 °C for 22 h. The  
precipitates of the reaction mixture were slurried in EtOAc/PE with chilling. The  
crude product was filtered, washed with EtOAc/PE, and dried *in vacuo* to afford the  
title compound as a purple solid (2.6 g). <sup>1</sup>H-NMR (CDCl<sub>3</sub>) δ: 2.25 (s, 6H, CH<sub>3</sub>), 2.45  
(s, 6H, CH<sub>3</sub>), 5.46 (d, 1H, *J* = 12.6 Hz, CH), 6.35 (s, 1H, pyrrolyl-H), 7.63 (d, 1H, *J* =  
10 12.6 Hz, CH).

**Example 2**

*[4-(2,4-Dimethyl-1H-pyrrol-3-yl)-pyrimidin-2-yl]-(4-fluoro-phenyl)-amine [1]*

To a mixture of 3-dimethylamino-1-(2,4-dimethyl-1H-pyrrol-3-yl)-propenone (1  
15 mmol, 0.19 g) and 4-fluorophenyl guanidine nitrate (2 mmol, 0.44 g) in 2-  
methoxyethanol (5 mL) was added NaOH (40 mg). The reaction mixture was heated  
at 100-120 °C under N<sub>2</sub> for 6 h. The solvent was evaporated to dryness and the residue  
was purified by flash chromatography (1:2 EtOAc/PE). Recrystallisation from  
EtOAc/PE afforded the title compound (174 mg, 62 %) as brown crystals. <sup>1</sup>H-NMR  
20 (CDCl<sub>3</sub>) δ: 2.21 (s, 3H, CH<sub>3</sub>), 2.43 (s, 3H, CH<sub>3</sub>), 6.33 (s, 1H, pyrrolyl-H), 6.73 (d, 1H,  
*J* = 5.3 Hz, pyrimidinyl-H), 7.00 (m, 2H, Ph-H), 7.79 (m, 2H, Ph-H), 8.28 (d, 1H, *J* =  
5.3 Hz, pyrimidinyl-H), 9.16 (s, 1H, NH), 10.59 (s, 1H, NH).

The following compounds were prepared in a manner analogous to that described  
25 above:

*[4-(2,4-Dimethyl-1H-pyrrol-3-yl)-pyrimidin-2-yl]-(3-nitro-phenyl)-amine [2]*

Yellow-orange solid. M.p. 197-199 °C. LC-MS:  $m/z$  = 310 (M+1).  $C_{16}H_{15}N_5O_2$  requires C, 62.12; H, 4.89; N, 22.64; found C, 62.61; H, 4.99; N, 22.20.  $^1H$ -NMR ( $CDCl_3$ )  $\delta$ : 2.71 (d, 6H,  $CH_3$ ), 7.05(d, 1H,  $J$  = 5.3 Hz, pyrimidinyl-H), 7.47 (m, 2H, Ph-H), 7.78 (m, 1H, Ph-H), 7.81 (s, 1H, Ar-H), 8.07 (m, 1H, Ph-H), 8.51 (d, 1H,  $J$  = 5.3 Hz, pyrimidinyl-H), 8.99 (br. s, 1H, NH), 9.91 (br. s, 1H, NH).

*[4-(2,4-Dimethyl-1H-pyrrol-3-yl)-pyrimidin-2-yl]-(4-iodo-phenyl)-amine [3]*

$^1H$ -NMR ( $CDCl_3$ )  $\delta$ : 2.26 (s, 3H,  $CH_3$ ), 2.48 (s, 3H,  $CH_3$ ), 6.80 (d, 1H,  $J$  = 5.3 Hz, pyrimidinyl-H), 7.47 (m, 2H, Ph-H), 7.57 (m, 2H, Ph-H), 7.23 (s, 1H, pyrrolyl-H), 8.32 (d, 1H,  $J$  = 5.3 Hz, pyrimidinyl-H).

*(3,4-Difluoro-phenyl)-[4-(2,4-dimethyl-1H-pyrrol-3-yl)-pyrimidin-2-yl]-amine [4]*

$^1H$ -NMR ( $CDCl_3$ )  $\delta$ : 2.23 (s, 3H,  $CH_3$ ), 2.46 (s, 3H,  $CH_3$ ), 6.42 (m, 1H, pyrrolyl-H), 6.77 (d, 1H,  $J$  = 5.3 Hz, pyrimidinyl-H), 7.11 (m, 1H, Ph-H), 7.41 (m, 1H, Ph-H), 8.05 (m, 1H, Ph-H), 8.29 (d, 1H,  $J$  = 5.3 Hz, pyrimidinyl-H), 9.21 (s, 1H, Ph-H), 10.46 (br. s, 1H, NH).

*(4-Chloro-phenyl)-[4-(2,4-dimethyl-1H-pyrrol-3-yl)-pyrimidin-2-yl]-amine [5]*

M.p. 219-223 °C. MS:  $[M+H]^+$  = 299.4 ( $C_{16}H_{15}ClN_4$  requires 298.8).  $^1H$ -NMR ( $DMSO-d_6$ )  $\delta$ : 2.19 (s, 3H,  $CH_3$ ), 2.42 (s, 3H,  $CH_3$ ), 6.48 (s, 1H, pyrrolyl-H), 6.82 (d, 1H,  $J$  = 5.3 Hz, pyrimidinyl-H), 7.31 (d, 2H,  $J$  = 8.7Hz, Ph-H), 7.84 (d, 2H,  $J$  = 8.7Hz, Ph-H), 8.34 (d, 1H,  $J$  = 5.3 Hz, pyrimidinyl-H), 9.45 (s, 1H, Ph-H), 10.72 (br. s, 1H, NH).

*(3,5-Difluoro-phenyl)-[4-(2,4-dimethyl-1H-pyrrol-3-yl)-pyrimidin-2-yl]-amine [6]*

M.p. 153.3-156.8 °C. MS:  $[M+H]^+$  = 303.6 ( $C_{16}H_{14}F_2N_4$  requires 300.3).  $^1H$ -NMR ( $CD_3OD$ )  $\delta$ : 2.25 (s, 3H,  $CH_3$ ), 2.47 (s, 3H,  $CH_3$ ), 6.42-6.48 (m, 2H, pyrrolyl-H and Ph-H), 6.80 (d, 1H,  $J$  = 5.5 Hz, pyrimidinyl-H), 7.44-7.48 (m, 2H, Ph-H), 8.31 (d, 1H,  $J$  = 5.5 Hz, pyrimidinyl-H).

*4-[4-(2,4-Dimethyl-1H-pyrrol-3-yl)-pyrimidin-2-ylamino]-phenol [7]*

M.p. 189.5-193.4 °C. MS:  $[M+H]^+ = 281.9$  ( $C_{16}H_{16}N_4O$  requires 280.3).  $^1H$ -NMR ( $CD_3OD$ )  $\delta$ : 2.23 (s, 3H,  $CH_3$ ), 2.43 (s, 3H,  $CH_3$ ), 6.44 (s, 1H, pyrrolyl-H), 6.75-6.78 (m, 3H, pyrimidinyl-H and Ph-H), 7.39 (d, 2H,  $J = 8.8$  Hz, Ph-H), 8.17 (d, 1H,  $J = 5.3$  Hz, pyrimidinyl-H).

*3-[4-(2,4-Dimethyl-1H-pyrrol-3-yl)-pyrimidin-2-ylamino]-phenol [8]*

M.p. 169.0-174.6 °C. MS:  $[M+H]^+ = 281.3$  ( $C_{16}H_{16}N_4O$  requires 280.3).  $^1H$ -NMR ( $CD_3OD$ )  $\delta$ : 2.26 (s, 3H,  $CH_3$ ), 2.47 (s, 3H,  $CH_3$ ), 6.44-6.48 (m, 2H, pyrrolyl-H, Ph-H), 6.84 (d, 1H,  $J = 5.4$  Hz, pyrimidinyl-H), 7.05-7.10 (m, 2H, Ph-H), 7.32 (m, 1H, Ph-H), 8.25 (d, 1H,  $J = 5.3$  Hz, pyrimidinyl-H).

*(2,4-Difluoro-phenyl)-[4-(2,4-dimethyl-1H-pyrrol-3-yl)-pyrimidin-2-yl]-amine [9]*

M.p. 219-220 °C. MS:  $[M+H]^+ = 302.6$  ( $C_{16}H_{14}F_2N_2$  requires 300.3).  $^1H$ -NMR ( $DMSO-d_6$ )  $\delta$ : 2.10 (s, 3H,  $CH_3$ ), 2.36 (s, 3H,  $CH_3$ ), 6.43 (s, 1H, pyrrolyl-H), 6.77 (d, 1H,  $J = 5.3$  Hz, pyrimidinyl-H), 7.06 (m, 1H, Ph-H), 7.27 (m, 1H, Ph-H), 7.66 (m, 1H, Ph-H), 8.25 (d, 1H,  $J = 5.3$  Hz, pyrimidinyl-H), 8.71 (s, 1H, Ph-H), 10.70 (br. s, 1H, NH).

*(2,4-Dichloro-phenyl)-[4-(2,4-dimethyl-1H-pyrrol-3-yl)-pyrimidin-2-yl]-amine [10]*

M.p. 158.5-159.7 °C. MS:  $[M+H]^+ = 335.4$  ( $C_{16}H_{14}Cl_2N_4$  requires 333.2).  $^1H$ -NMR ( $DMSO-d_6$ )  $\delta$ : 2.18 (s, 3H,  $CH_3$ ), 2.38 (s, 3H,  $CH_3$ ), 6.51 (s, 1H, pyrrolyl-H), 6.90 (d, 1H,  $J = 5.5$  Hz, pyrimidinyl-H), 7.46 (m, 1H, Ph-H), 7.71 (m, 1H, Ph-H), 8.05 (m, 1H, Ph-H), 8.36 (d, 1H,  $J = 5.3$  Hz, pyrimidinyl-H), 8.49 (s, 1H, Ph-H), 10.80 (br. s, 1H, NH).

*(4-Chloro-3-trifluoromethyl-phenyl)-[4-(2,4-dimethyl-1H-pyrrol-3-yl)-pyrimidin-2-yl]-amine [11]*

M.p. 187.7-190.7 °C. MS:  $[M+H]^+ = 368.6$  ( $C_{17}H_{14}ClF_3N_4$  requires 366.8).  $^1H$ -NMR (DMSO- $d_6$ )  $\delta$ : 2.19 (s, 3H,  $CH_3$ ), 2.42 (s, 3H,  $CH_3$ ), 6.50 (s, 1H, pyrrolyl-H), 6.89 (d, 1H,  $J = 5.3$  Hz, pyrimidinyl-H), 7.61 (m, 1H, Ph-H), 8.08 (m, 1H, Ph-H), 8.39-8.42 (m, 2H, Ph-H and pyrimidinyl-H), 9.79 (s, 1H), 10.80 (br. s, 1H, NH).

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*[4-(2,4-Dimethyl-1H-pyrrol-3-yl)-pyrimidin-2-yl]-(4-trifluoromethyl-phenyl)-amine*  
**[12]**

M.p. 165.6-167.9 °C. MS:  $[M+H]^+ = 332.9$  ( $C_{17}H_{15}F_3N_4$  requires 332.3).  $^1H$ -NMR (DMSO- $d_6$ )  $\delta$ : 2.26 (s, 3H,  $CH_3$ ), 2.49 (s, 3H,  $CH_3$ ), 6.56 (s, 1H, pyrrolyl-H), 6.94 (d, 1H,  $J = 5.3$  Hz, pyrimidinyl-H), 7.67 (d, 2H,  $J = 8.5$  Hz, Ph-H), 8.09 (d, 2H,  $J = 8.5$  Hz, Ph-H), 8.45 (d, 1H,  $J = 5.3$  Hz, pyrimidinyl-H), 9.82 (s, 1H, NH).

10

*[4-(2,4-Dimethyl-1H-pyrrol-3-yl)-pyrimidin-2-yl]-(3-trifluoromethyl-phenyl)-amine*  
**[13]**

M.p. 152.7-154.3 °C. MS:  $[M+H]^+ = 332.6$  ( $C_{17}H_{15}F_3N_4$  requires 332.3).  $^1H$ -NMR (DMSO- $d_6$ )  $\delta$ : 2.26 (s, 3H,  $CH_3$ ), 2.48 (s, 3H,  $CH_3$ ), 6.56 (s, 1H, pyrrolyl-H), 6.92 (d, 1H,  $J = 5.3$  Hz, pyrimidinyl-H), 7.29 (m, 1H, Ph-H), 7.55 (m, 1H, Ph-H), 8.03 (m, 1H, Ph-H), 8.42-7.45 (m, 2H, pyrimidinyl-H and Ph-H), 9.73 (s, 1H, NH), 10.83 (br. s, 1H, NH).

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*(3-Chloro-phenyl)-[4-(2,4-dimethyl-1H-pyrrol-3-yl)-pyrimidin-2-yl]-amine* **[14]**

M.p. 140.4-144.2 °C. MS:  $[M+H]^+ = 299.5$  ( $C_{16}H_{15}ClN_4$  requires 298.8).  $^1H$ -NMR (DMSO- $d_6$ )  $\delta$ : 2.21 (s, 3H,  $CH_3$ ), 2.44 (s, 3H,  $CH_3$ ), 6.51 (s, 1H, pyrrolyl-H), 6.85 (d, 1H,  $J = 5.3$  Hz, pyrimidinyl-H), 6.94 (d, 1H,  $J = 7.6$  Hz, Ph-H), 7.28 (t, 1H,  $J = 8.1$  Hz, Ph-H), 7.61 (d, 1H,  $J = 8.2$  Hz, Ph-H), 8.19 (s, 1H, Ph-H), 8.37 (d, 1H,  $J = 5.3$  Hz, pyrimidinyl-H), 9.55 (s, 1H, NH), 10.78 (br. s, 1H, NH).

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*N-[4-(2,4-Dimethyl-1H-pyrrol-3-yl)-pyrimidin-2-yl]-N',N'-dimethyl-benzene-1,4-diamine* **[15]**

M.p. 179.9-182.1 °C. MS:  $[M+H]^+ = 307.3$  ( $C_{18}H_{21}N_5$  requires 307.4).  $^1H$ -NMR ( $CDCl_3$ )  $\delta$ : 2.25 (s, 3H,  $CH_3$ ), 2.46 (s, 3H,  $CH_3$ ), 2.91 (s, 6H,  $CH_3$ ), 6.46 (s, 1H, pyrrolyl-H), 6.70 (d, 1H,  $J = 5.1$  Hz, pyrimidinyl-H), 6.78 (dd, 2H,  $J = 6.8, 2.2$  Hz, Ph-H), 6.79 (br. s, 1H, NH), 7.45 (dd, 2H,  $J = 6.8, 2.2$  Hz, Ph-H), 7.80 (br. s, 1H, NH), 8.28 (d, 1H,  $J = 5.1$  Hz, pyrimidinyl-H).

*(3-Chloro-4-iodo-phenyl)-[4-(2,4-dimethyl-1H-pyrrol-3-yl)-pyrimidin-2-yl]-amine*

**[16]**

M.p. 185.0-187.4 °C. MS:  $[M+H]^+ = 423.9$  ( $C_{16}H_{14}ClIN_4$  requires 424.7).  $^1H$ -NMR ( $DMSO-d_6$ )  $\delta$ : 2.18 (s, 3H,  $CH_3$ ), 2.41 (s, 3H,  $CH_3$ ), 6.48 (s, 1H, pyrrolyl-H), 6.84 (d, 1H,  $J = 5.4$  Hz, pyrimidinyl-H), 7.40 (dd, 1H,  $J = 8.8, 2.4$  Hz, Ph-H), 7.75 (d, 1H,  $J = 8.8$  Hz, Ph-H), 8.34 (m, 1H, Ph-H), 8.36 (d, 1H,  $J = 5.4$  Hz, pyrimidinyl-H), 9.61 (s, 1H, NH), 10.75 (s, 1H, NH).

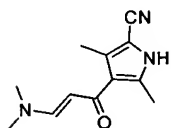
*[4-(2,4-Dimethyl-1H-pyrrol-3-yl)-pyrimidin-2-yl]-(3-fluoro-4-iodo-phenyl)-amine*

**[17]**

M.p. 200-202 °C. MS:  $[M+H]^+ = 407.4$  ( $C_{16}H_{14}FIN_4$  requires 408.2).  $^1H$ -NMR ( $DMSO-d_6$ )  $\delta$ : 2.18 (s, 3H,  $CH_3$ ), 2.40 (s, 3H,  $CH_3$ ), 6.48 (s, 1H, pyrrolyl-H), 6.84 (d, 1H,  $J = 5.4$  Hz, pyrimidinyl-H), 7.35 (m, 1H, Ph-H), 7.64 (t, 1H,  $J = 8.0$  Hz, Ph-H), 8.02 (dd, 1H,  $J = 12.0, 2.2$  Hz, Ph-H), 8.36 (d, 1H,  $J = 5.4$  Hz, pyrimidinyl-H), 9.65 (s, 1H, NH), 10.75 (s, 1H, NH).

**Example 3**

*4-(3-Dimethylamino-acryloyl)-3,5-dimethyl-1H-pyrrole-2-carbonitrile*



Ethyl cyanoacetate (10 mL, 94 mmol) was diluted with AcOH (20 mL) and the solution was cooled to  $-10$  °C (ice-MeOH bath).  $NaNO_2$  (6.5 g, 94 mmol) was



dissolved in H<sub>2</sub>O (10 mL) and the solution was added dropwise over a period of 40 min, keeping the internal temperature < 0° C. After completion of the addition, the reaction mixture was stirred for 1h with cooling. It was then warmed to room temperature and stirred for a further 3 h. The mixture was diluted with acetic acid (50 mL) and H<sub>2</sub>O (50 mL). Pentane-2,4-dione (10.6 mL, 103 mmol) was added and the mixture heated to ~75 °C. To this reaction mixture Zn powder (6.9 g, 105 mmol) was added in portions over a period of 30 min at such a rate as to maintain the internal temperature < 90 °C. The reaction mixture was then heated for a further 30 min before pouring into H<sub>2</sub>O (1 mL). From the reaction mixture 3,5-dimethyl-1*H*-pyrrole-2-carbonitrile (3.67 g) was filtered as an off-white solid. The filtrate was extracted with EtOAc (3 × 500 mL). The combined organic extracts were washed (brine) and dried (MgSO<sub>4</sub>). The solvent was evaporated to a brown oil, which was purified by chromatography (100 g SiO<sub>2</sub>, eluted with 4:1 heptane / EtOAc) to afford a further crop (4.41 g) of this product as a pale yellow solid (total yield 72 %).

3,5-Dimethyl-1*H*-pyrrole-2-carbonitrile (1.2 g, 10 mmol) was dissolved in anhydrous 1,2-dichloroethane (15 mL) and AlCl<sub>3</sub> (2.93 g, 22 mmol) was added proportion-wise. The reaction vessel was purged with N<sub>2</sub> and was cooled in an ice-water bath. AcCl (0.71 mL, 10 mmol) was added dropwise and the mixture was stirred for 1 h with cooling and for a further 3 h at room temperature. The reaction mixture was quenched by careful addition of 2 M aq HCl. The acidity of the mixture was adjusted to approximately pH 6 by addition of NaHCO<sub>3</sub>. After separation of the organic phase, the aqueous phase was extracted with EtOAc (3 × 100mL). The combined organic phases were washed (H<sub>2</sub>O, then brine), dried (MgSO<sub>4</sub>), and filtered. The solvent was evaporated to afford of 4-acetyl-3,5-dimethyl-1*H*-pyrrole-2-carbonitrile (1.42 g, 88 %) as a pale tan solid. <sup>1</sup>H-NMR (CDCl<sub>3</sub>) δ: 2.44 (s, 3H, CH<sub>3</sub>), 2.45 (s, 3H, CH<sub>3</sub>), 2.54 (s, 3H, CH<sub>3</sub>), 8.75 (br. s, 1H, NH).

4-Acetyl-3,5-dimethyl-1*H*-pyrrole-2-carbonitrile (1.38 g, 8.51 mmol) was suspended in 1,1-bis-dimethylamino-3,3-dimethyl-butan-2-one (1.3 mL) and heated at 75 °C for 42 h. The reaction mixture was evaporated to dryness and the residue was

purified by SiO<sub>2</sub> chromatography (heptane / EtOAc) to afford the title compound (1.2 g, 65 %) as a pale tan solid. <sup>1</sup>H-NMR (DMSO-d<sub>6</sub>) δ: 2.21 (s, 3H, CH<sub>3</sub>), 2.31 (s, 3H, CH<sub>3</sub>), 3.32 (s, 6H, CH<sub>3</sub>), 5.22 (d, 1H, *J* = 12.4 Hz, CH), 7.47 (d 1H, *J* = 12.4 Hz, CH), 11.96 (br. s, 1H, NH).

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Example 4

*3,5-Dimethyl-4-[2-(3-nitro-phenylamino)-pyrimidin-4-yl]-1H-pyrrole-2-carbonitrile*  
**[31]**

To a mixture of 4-(3-dimethylamino-acryloyl)-3,5-dimethyl-1H-pyrrole-2-carbonitrile  
10 (1.0 mmol, 0.22 g) and 3-nitrophenyl guanidine nitrate (1.5 mmol, 0.36 g) in 2-methoxyethanol (5 mL) was added K<sub>2</sub>CO<sub>3</sub> (138 mg, 1.0 mmol). The reaction mixture was heated at 120 °C under N<sub>2</sub> for 18 h. The solvent was evaporated to dryness and the residue was purified by flash chromatography (1:2 EtOAc / heptane) to afford the title compound as a light-yellow solid. M.p. 258-259 °C. MS: [M+H]<sup>+</sup> = 336.1  
15 (C<sub>17</sub>H<sub>14</sub>N<sub>6</sub>O<sub>2</sub> requires 334.3). <sup>1</sup>H-NMR (CD<sub>3</sub>OD) δ: 2.39 (s, 3H, CH<sub>3</sub>), 2.49 (s, 3H, CH<sub>3</sub>), 6.94 (d, 1H, *J* = 5.1 Hz, pyrimidinyl-H), 7.50 (t, 1H, *J* = 8.3 Hz, Ph-H), 7.81 (m, 1H, Ph-H), 7.94 (m, 1H, Ph-H), 8.45 (d, 1H, *J* = 5.1 Hz, pyrimidinyl-H), 8.94 (t, 1H, *J* = 2.2 Hz, Ph-H).

20 The following compounds were prepared in a manner analogous to that described above:

*4-[2-(4-Fluoro-phenylamino)-pyrimidin-4-yl]-3,5-dimethyl-1H-pyrrole-2-carbonitrile*  
**[32]**

25 MS: [M+H]<sup>+</sup> = 307.7 (C<sub>17</sub>H<sub>14</sub>FN<sub>5</sub> requires 307.3). <sup>1</sup>H-NMR (DMSO-d<sub>6</sub>) δ: 2.30 (s, 3H, CH<sub>3</sub>), 2.40 (s, 3H, CH<sub>3</sub>), 6.84 (d, 1H, *J* = 5.0 Hz, pyrimidinyl -H), 7.00 (m, 2H, Ph-H), 7.73 (m, 2H, Ph-H), 8.40 (d, 1H, *J* = 5.5 Hz, pyrimidinyl -H), 9.46 (s, 1H, NH), 12.19 (br. s, 1H, NH).

*4-[2-(4-Hydroxy-phenylamino)-pyrimidin-4-yl]-3,5-dimethyl-1H-pyrrole-2-carbonitrile [33]*

M.p. 272-276 °C. MS:  $[M+H]^+ = 305.8$  ( $C_{17}H_{15}N_5O$  requires 305.3).  $^1H$ -NMR (CD<sub>3</sub>OD)  $\delta$ : 2.33 (s, 3H, CH<sub>3</sub>), 2.41 (s, 3H, CH<sub>3</sub>), 6.74-6.56 (m, 3H, pyrimidinyl-H/Ph-H), 7.36 (d, 2H,  $J = 8.5$  Hz, Ph-H), 8.25 (d, 1H,  $J = 5.4$  Hz, pyrimidinyl-H).

*3,5-Dimethyl-4-[2-(4-trifluoromethyl-phenylamino)-pyrimidin-4-yl]-1H-pyrrole-2-carbonitrile [34]*

M.p. 195.6-198.9 °C. MS:  $[M+H]^+ = 357.7$  ( $C_{18}H_{14}F_3N_5$  requires 357.3).  $^1H$ -NMR (CDCl<sub>3</sub>)  $\delta$ : 2.33 (s, 3H, CH<sub>3</sub>), 2.44 (s, 3H, CH<sub>3</sub>), 6.75 (d, 1H,  $J = 5.1$  Hz, pyrimidinyl-H), 7.20 (br. s, 1H, NH), 7.50 (d, 2H,  $J = 8.8$  Hz, Ph-H), 7.71 (d, 2H,  $J = 8.8$  Hz, Ph-H), 8.39 (d, 1H,  $J = 5.1$  Hz, pyrimidinyl), 8.40 (br. s, 1H, NH).

*4-[2-(4-Iodo-phenylamino)-pyrimidin-4-yl]-3,5-dimethyl-1H-pyrrole-2-carbonitrile [35]*

M.p. 178.3-181.2 °C. MS:  $[M+H]^+ = 416.6$  ( $C_{18}H_{14}IN_5$  requires 415.2).  $^1H$ -NMR (CDCl<sub>3</sub>)  $\delta$ : 2.39 (s, 3H, CH<sub>3</sub>), 2.49 (s, 3H, CH<sub>3</sub>), 6.76 (d, 1H,  $J = 5.1$  Hz, pyrimidinyl-H), 7.10 (br. s, 1H, NH), 7.44 (d, 2H,  $J = 8.8$  Hz, Ph-H), 7.61 (d, 2H,  $J = 8.8$  Hz, Ph-H), 8.42 (d, 1H,  $J = 5.1$  Hz, pyrimidinyl-H), 8.45 (br. s, 1H, NH).

*4-[2-(3-Hydroxy-phenylamino)-pyrimidin-4-yl]-3,5-dimethyl-1H-pyrrole-2-carbonitrile [36]*

M.p. 247-250 °C. MS:  $[M+H]^+ = 305.8$  ( $C_{17}H_{15}N_5O$  requires 305.3).  $^1H$ -NMR (DMSO-d<sub>6</sub>)  $\delta$ : 2.31 (s, 3H, CH<sub>3</sub>), 2.41 (s, 3H, CH<sub>3</sub>), 6.33 (m, 1H, Ph-H), 6.82 (d, 1H,  $J = 5.1$  Hz, pyrimidinyl-H), 7.01 (t, 1H,  $J = 8.1$  Hz, Ph-H), 7.11 (m, 1H, Ph-H), 7.33 (t, 1H,  $J = 2.1$  Hz, Ph-H), 8.40 (d, 1H,  $J = 5.4$  Hz, pyrimidinyl-H), 9.18 (s, 1H), 9.30 (s, 1H), 12.20 (br. s, 1H, NH).

3,5-Dimethyl-4-[2-(4-methyl-3-nitro-phenylamino)-pyrimidin-4-yl]-1H-pyrrole-2-carbonitrile [37]

M.p. 233-237 °C. MS:  $[M+H]^+ = 350.0$  ( $C_{18}H_{16}N_6O_2$  requires 348.6).  $^1H$ -NMR (DMSO- $d_6$ )  $\delta$ : 2.31 (s, 3H, CH<sub>3</sub>), 2.42 (s, 3H, CH<sub>3</sub>), 2.44 (s, 3H, CH<sub>3</sub>), 6.92 (d, 1H,  $J = 5.4$  Hz, pyrimidinyl-H), 7.39 (d, 1H,  $J = 8.5$  Hz, Ph-H), 7.87 (dd, 1H,  $J = 8.1, 1.7$  Hz, Ph-H), 8.48 (d, 1H,  $J = 5.1$  Hz, pyrimidinyl-H), 8.63 (d, 1H,  $J = 1.7$  Hz, Ph-H), 9.87 (s, 1H, NH), 12.21 (br. s, 1H, NH).

4-[2-(3-Iodo-4-methyl-phenylamino)-pyrimidin-4-yl]-3,5-dimethyl-1H-pyrrole-2-carbonitrile [38]

M.p. 189.5-191.7 °C. MS:  $[M+H]^+ = 431.5$  ( $C_{18}H_{16}IN_5$  requires 429.6).  $^1H$ -NMR (DMSO- $d_6$ )  $\delta$ : 2.29 (s, 3H, CH<sub>3</sub>), 2.32 (s, 3H, CH<sub>3</sub>), 2.43 (s, 3H, CH<sub>3</sub>), 6.85 (d, 1H,  $J = 5.1$  Hz, pyrimidinyl-H), 7.20 (d, 1H,  $J = 8.1$  Hz, Ph-H), 7.57 (m, 1H, Ph-H), 8.41-8.43 (m, 2H, Ph-H, pyrimidinyl-H), 9.48 (s, 1H, NH), 12.20 (br. s, 1H, NH).

4-[2-(4-Chloro-3-methyl-phenylamino)-pyrimidin-4-yl]-3,5-dimethyl-1H-pyrrole-2-carbonitrile [39]

M.p. 194.2-197.9 °C. MS:  $[M+H]^+ = 338.0$  ( $C_{18}H_{16}ClN_5$  requires 337.8).  $^1H$ -NMR (DMSO- $d_6$ )  $\delta$ : 2.27 (s, 3H, CH<sub>3</sub>), 2.31 (s, 3H, CH<sub>3</sub>), 2.41 (s, 3H, CH<sub>3</sub>), 6.86 (d, 1H,  $J = 5.4$  Hz, pyrimidinyl-H), 7.28 (d, 1H,  $J = 8.5$  Hz, Ph-H), 7.61 (dd, 1H,  $J = 8.8, 2.4$  Hz, Ph-H), 7.73 (d, 1H,  $J = 2.7$  Hz, Ph-H), 8.43 (d, 1H,  $J = 5.1$  Hz, pyrimidinyl-H), 9.51 (s, 1H, NH), 12.21 (br. s, 1H, NH).

4-[2-(3-Hydroxy-4-methyl-phenylamino)-pyrimidin-4-yl]-3,5-dimethyl-1H-pyrrole-2-carbonitrile [40]

M.p. 221-225 °C. MS:  $[M+H]^+ = 320.9$  ( $C_{18}H_{17}N_5O$  requires 319.4).  $^1H$ -NMR (DMSO- $d_6$ )  $\delta$ : 2.03 (s, 3H, CH<sub>3</sub>), 2.29 (s, 3H, CH<sub>3</sub>), 2.40 (s, 3H, CH<sub>3</sub>), 6.78 (d, 1H,  $J = 5.1$  Hz, pyrimidinyl-H), 6.89 (d, 1H,  $J = 8.1$  Hz, Ph-H), 7.02 (dd, 1H,  $J = 8.3,$

1.7 Hz, *Ph-H*), 7.29 (*d*, 1H, *J* = 0.7 Hz, *Ph-H*), 8.37 (*d*, 1H, *J* = 4.9 Hz, *pyrimidinyl-H*), 9.08 (*s*, 1H), 9.20 (*s*, 1H), 12.17 (*br. s*, 1H, *NH*).

4-[2-(4-Fluoro-3-methyl-phenylamino)-pyrimidin-4-yl]-3,5-dimethyl-1H-pyrrole-2-carbonitrile **[41]**

*M.p.* 161.3-164.1 °C. *MS*:  $[M+H]^+ = 321.6$  ( $C_{18}H_{16}FN_5$  requires 321.4).  $^1H$ -NMR ( $DMSO-d_6$ )  $\delta$ : 2.19 (*s*, 3H,  $CH_3$ ), 2.30 (*s*, 3H,  $CH_3$ ), 2.42 (*s*, 3H,  $CH_3$ ), 6.82 (*d*, 1H, *J* = 5.1 Hz, *pyrimidinyl-H*), 7.03 (*t*, 1H, *J* = 9.3 Hz, *Ph-H*), 7.53 (*m*, 1H, *Ph-H*), 7.61 (*dd*, 1H, *J* = 7.1, 2.4 Hz, *Ph-H*), 8.39 (*d*, 1H, *J* = 5.1 Hz, *pyrimidinyl-H*), 9.36 (*s*, 1H, *NH*), 12.20 (*br. s*, 1H, *NH*).

4-[2-(3-Fluoro-4-methyl-phenylamino)-pyrimidin-4-yl]-3,5-dimethyl-1H-pyrrole-2-carbonitrile **[42]**

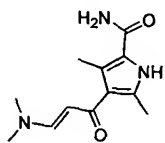
*M.p.* 177.7-179.9 °C. *MS*:  $[M+H]^+ = 322.5$  ( $C_{18}H_{16}FN_5$  requires 321.3).  $^1H$ -NMR ( $DMSO-d_6$ )  $\delta$ : 2.15 (*s*, 3H,  $CH_3$ ), 2.30 (*s*, 3H,  $CH_3$ ), 2.43 (*s*, 3H,  $CH_3$ ), 6.86 (*d*, 1H, *J* = 5.1 Hz, *pyrimidinyl-H*), 7.13 (*t*, 1H, *J* = 9.0 Hz, *Ph-H*), 7.36 (*dd*, 1H, *J* = 8.1, 1.7 Hz, *Ph-H*), 7.75 (*dd*, 1H, *J* = 12.9, 1.5 Hz, *Ph-H*), 8.43 (*d*, 1H, *J* = 5.4 Hz, *pyrimidinyl-H*), 9.56 (*s*, 1H, *NH*), 12.21 (*br. s*, 1H, *NH*).

4-[2-(4-Dimethylamino-phenylamino)-pyrimidin-4-yl]-3,5-dimethyl-1H-pyrrole-2-carbonitrile **[43]**

*M.p.* 190.6-193.7 °C. *MS*:  $[M+H]^+ = 334.7$  ( $C_{19}H_{20}N_6$  requires 332.4).  $^1H$ -NMR ( $CDCl_3$ )  $\delta$ : 2.36 (*s*, 3H,  $CH_3$ ), 2.46 (*s*, 3H,  $CH_3$ ), 2.94 (*br. s*, 6H,  $CH_3$ ), 6.66 (*d*, 1H, *J* = 5.6 Hz, *pyrimidinyl-H*), 6.79-6.80 (*m*, 2H, *Ph-H*), 7.05 (*br. s*, 1H, *NH*), 7.40-7.43 (*m*, 2H, *Ph-H*), 8.34 (*d*, 1H, *J* = 5.1 Hz, *pyrimidinyl-H*), 8.52 (*br. s*, 1H, *NH*).

#### Example 5

4-(3-Dimethylamino-acryloyl)-3,5-dimethyl-1H-pyrrole-2-carboxylic acid amide



1-(2,4-Dimethyl-1*H*-pyrrol-3-yl)-ethanone (1.1 g, 10 mmol) was partially dissolved in 2 M solution of ammonia in MeOH and H<sub>2</sub>O<sub>2</sub> (10 mL of a 27 % w/w solution in H<sub>2</sub>O) was added dropwise over a period of 40 min at such a rate as to maintain the internal temperature  $\leq 30$  °C. The mixture was stirred for 18 h at room temperature. The resulting suspended white solid was filtered and recrystallised from EtOAc to afford 4-acetyl-3,5-dimethyl-1*H*-pyrrole-2-carboxylic acid amide (1.06 g). An aliquot (720 mg, 4 mmol) was suspended in 1,1-bis-dimethylamino-3,3-dimethyl-butan-2-one (2mL, 9.6mmol) in a N<sub>2</sub>-flushed flask and heated at 75 °C for 48 h. The crude mixture was cooled and purified by SiO<sub>2</sub> chromatography (EtOAc / MeOH gradient elution). The title compound (449 mg) was obtained as a buff solid. <sup>1</sup>H-NMR (DMSO-d<sub>6</sub>)  $\delta$ : 2.30 (s, 3H, CH<sub>3</sub>), 2.46 (s, 3H, CH<sub>3</sub>), 2.90 (br. s, 2H, NH). 3.09 (s, 3H, CH<sub>3</sub>), 3.13 (s, 3H, CH<sub>3</sub>), 5.23 (d, 1H, *J* = 12.4 Hz, CH), 7.38 (d, 1H, *J* = 12.7 Hz, CH), 10.97 (br. s, 1H, NH).

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#### Example 6

*4-[2-(4-Fluoro-phenylamino)-pyrimidin-4-yl]-3,5-dimethyl-1H-pyrrole-2-carboxylic acid amide [44]*

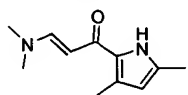
4-(3-Dimethylamino-acryloyl)-3,5-dimethyl-1*H*-pyrrole-2-carboxylic acid amide (100 mg, 0.43 mmol), 4-fluorophenylguanidine nitrate (139 mg, 0.65 mmol) and K<sub>2</sub>CO<sub>3</sub> (94 mg, 0.68 mmol) were partially dissolved in 2-methoxyethanol (5 mL) and heated at 120 °C for 18 h. The mixture was concentrated *in vacuo* and purified by SiO<sub>2</sub> chromatography (EtOAc / MeOH gradient elution). The crude product was triturated in iPr<sub>2</sub>O to afford the title compound (31 mg) as a buff solid. M.p. 93.5-96.8 °C. MS: [M+H<sup>+</sup>] = 326.9 (C<sub>17</sub>H<sub>16</sub>FN<sub>5</sub>O requires 325.3). <sup>1</sup>H-NMR (DMSO-d<sub>6</sub>)  $\delta$ : 2.36 (s, 3H, CH<sub>3</sub>), 2.39 (s, 3H, CH<sub>3</sub>), 6.79 (d, 1H, *J* = 5.4 Hz, pyrimidinyl-H), 6.92 (br. s, 2H,

25

NH), 7.07 (t, 2H,  $J = 8.5$  Hz, Ph-H), 7.76-7.78 (m, 2H, Ph-H), 8.36 (d, 1H,  $J = 5.4$  Hz, pyrimidinyl-H), 9.41 (s, 1H, NH), 11.24 (br. s, 1H, NH).

### Example 7

#### 5    3-Dimethylamino-1-(3,5-dimethyl-1H-pyrrol-2-yl)-propenone



Pentane-2,4-dione (10.3 mL, 0.1 mol) was diluted with AcOH and cooled to 0 °C. NaNO<sub>2</sub> (6.9 g, 0.1 mol) was dissolved in H<sub>2</sub>O (10 mL) and added dropwise, keeping the internal temperature ≤ 10 °C. The mixture was stirred for 1 h with cooling then 3 h  
10 at room temperature. Ethyl acetoacetate (14 mL, 0.11 mol) was dissolved in 1:1 AcOH / H<sub>2</sub>O (100 mL) and Zn powder (7.19 g, 0.11 mol) was added. To this the oxime solution was added and the mixture was heated at 100 °C for 30 min then poured into H<sub>2</sub>O (0.8 L). The aqueous mixture was extracted with EtOAc (3 × 500mL). The combined organic extracts were washed (brine), dried (MgSO<sub>4</sub>), filtered,  
15 and evaporated *in vacuo*. The residue was purified by SiO<sub>2</sub> chromatography (heptane / EtOAc gradient elution) to afford 5-acetyl-2,4-dimethyl-1H-pyrrole-3-carboxylic acid ethyl ester (4.32 g). An aliquot (3.5 g, 16.7 mmol) was added to KOH (4.22 g, 75.3 mmol) dissolved in H<sub>2</sub>O (5 mL) and the mixture was heated at 130 °C for 18 h. It was then cooled, diluted with H<sub>2</sub>O (50 mL) and acidified with 2 M aq HCl. This mixture  
20 was extracted with EtOAc (3 × 50 mL). The combined organic extracts were washed (brine), dried (MgSO<sub>4</sub>), filtered, and evaporated *in vacuo*. The residue was purified by SiO<sub>2</sub> chromatography (EtOAc) to afford 1-(3,5-dimethyl-1H-pyrrol-2-yl)-ethanone (1.05 g). An aliquot (536 mg, 3.9 mmol) was suspended in *N,N*-dimethylformamide dimethyl acetal (1.2 mL, 8.8 mmol) in a N<sub>2</sub>-flushed flask and heated at 90 °C for 48 h.  
25 The mixture was cooled and triturated in cold EtOAc. The resulting precipitate was filtered and washed with EtOAc to afford the title compound (444 mg) as a buff. <sup>1</sup>H-NMR (DMSO-d<sub>6</sub>) δ: 1.41 (s, 3H, CH<sub>3</sub>), 2.23 (s, 3H, CH<sub>3</sub>), 2.23 (br. s, 6H, CH<sub>3</sub>), 4.77 (d, 1H,  $J = 12.4$  Hz, CH), 4.95 (s, 1H, pyrrolyl-H), 6.85 (d, 1H,  $J = 12.4$  Hz, CH).

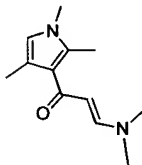
Example 8

*[4-(3,5-Dimethyl-1H-pyrrol-2-yl)-pyrimidin-2-yl]-(4-fluoro-phenyl)-amine* [45]

3-Dimethylamino-1-(3,5-dimethyl-1H-pyrrol-2-yl)-propenone (125 mg, 0.65 mmol),  
 5 4-fluorophenyl guanidine nitrate (211 mg, 0.98 mmol), and K<sub>2</sub>CO<sub>3</sub> (149 mg, 1.08 mmol) were partially dissolved in 2-methoxyethanol and heated at 120 °C for 18 h. The mixture was concentrated *in vacuo* and purified by SiO<sub>2</sub> chromatography (5-g Isolute SI™ cartridge eluted with an heptane / EtOAc gradient). The crude product was triturated in iPr<sub>2</sub>O to afford the title compound (158 mg) as a buff solid. M.p. 168.4-  
 10 171.5 °C. MS: [M+H]<sup>+</sup> = 283.9 (C<sub>16</sub>H<sub>15</sub>FN<sub>4</sub> requires 282.3). <sup>1</sup>H-NMR (CDCl<sub>3</sub>) δ: 2.30 (s, 3H, CH<sub>3</sub>), 2.35 (s, 3H, CH<sub>3</sub>), 5.85 (s, 1H, pyrrolyl-H), 6.83 (d, 1H, *J* = 5.6 Hz, pyrimidinyl-H), 6.87 (br. s, 1H, NH), 7.05 (t, 2H, *J* = 8.5 Hz, Ph-H), 7.51-7.54 (m, 2H, Ph-H), 8.26 (d, 1H, *J* = 5.4 Hz, pyrimidinyl-H), 9.08 (br. s, 1H, NH).

15 Example 9

*3-Dimethylamino-1-(1,2,4-trimethyl-1H-pyrrol-3-yl)-propenone*



KOH (818 mg, 14.6 mmol) was partially dissolved in DMSO (15 mL) and stirred for 5 min. 1-(2,4-dimethyl-1H-pyrrol-3-yl)-ethanone (1 g, 7.3 mmol) was added in small  
 20 portions and the mixture was stirred for 45 min. Iodomethane (0.54 mL, 8.75 mmol) was added dropwise at such a rate as to maintain the internal temperature ≤ 30 °C. The mixture was stirred for a further 45 min then poured into H<sub>2</sub>O (50 mL) and extracted with Et<sub>2</sub>O (3 × 60 mL). The combined organic extracts were washed (brine), dried (MgSO<sub>4</sub>), filtered, and evaporated *in vacuo* to afford 1-(1,2,4-trimethyl-1H-pyrrol-3-yl)-ethanone (1.06 g) as a pink solid. This was suspended in 1,1-bis-dimethylamino-3,3-dimethyl-butan-2-one (3.6 mL, 17.5 mmol) in a N<sub>2</sub>-flushed flask and heated at 70  
 25 °C for 36 h. The mixture was evaporated *in vacuo* and the residue triturated in EtOAc.



The titled compound (973 mg) was obtained as a reddish solid.  $^1\text{H-NMR}$  ( $\text{DMSO-d}_6$ )  $\delta$ : 1.30 (s, 3H,  $\text{CH}_3$ ), 1.51 (s, 3H,  $\text{CH}_3$ ), 2.20 (br. s, 6H,  $\text{CH}_3$ ), 2.66 (s, 3H,  $\text{CH}_3$ ), 4.57 (d, 1H,  $J = 12.5$  Hz, CH), 5.53 (s, 1H, pyrrolyl-H), 6.72 (d, 1H,  $J = 12.7$  Hz, CH).

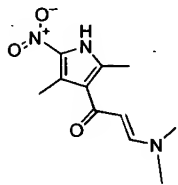
5 Example 10

*(4-Fluoro-phenyl)-[4-(1,2,4-trimethyl-1H-pyrrol-3-yl)-pyrimidin-2-yl]-amine [46]*

3-Dimethylamino-1-(1,2,4-trimethyl-1H-pyrrol-3-yl)-propenone (150 mg, 0.73 mmol), 4-fluorophenyl guanidine nitrate (189 mg, 0.87 mmol) and  $\text{K}_2\text{CO}_3$  (144 mg, 1.04 mmol) were partially dissolved in 2-methoxyethanol (3 mL) and heated at 110 °C  
10 36 h. The mixture was concentrated in *vacuo* and purified by  $\text{SiO}_2$  chromatography (heptane / EtOAc gradient elution). The title compound (20 mg) was obtained as a brownish solid after recrystallisation from  $\text{iPr}_2\text{O}$  / heptane. M.p. 124.2-128.3 °C. MS:  $[\text{M}+\text{H}]^+ = 298.4$  ( $\text{C}_{17}\text{H}_{17}\text{FN}_4$  requires 296.3).  $^1\text{H-NMR}$  ( $\text{CDCl}_3$ )  $\delta$ : 2.21 (s, 3H,  $\text{CH}_3$ ), 2.42 (s, 3H,  $\text{CH}_3$ ), 3.51 (s, 3H,  $\text{CH}_3$ ), 6.40 (s, 1H, pyrrolyl-H), 6.76 (d, 1H,  $J = 5.4$  Hz, pyrimidinyl-H), 7.01 (t, 2H,  $J = 8.8$  Hz, Ph-H), 7.58-7.61 (m, 2H, Ph-H), 7.70 (br. s, 1H, NH), 8.28 (d, 1H,  $J = 5.6$  Hz, pyrimidinyl-H).

Example 11

*3-Dimethylamino-1-(2,4-dimethyl-5-nitro-1H-pyrrol-3-yl)-propenone*



$\text{HNO}_3$  (0.28 mL of a 69 % w/v aq solution, 4.37 mmol) was added dropwise to  $\text{Ac}_2\text{O}$  (5 mL) at room temperature, keeping the internal temperature  $\leq 25$  °C. The nitrating mixture was stirred at room temperature for 15 min before cooling to -40 °C. 1-(2,4-dimethyl-1H-pyrrol-3-yl)-ethanone (500 mg, 3.64 mmol) was dissolved in  $\text{Ac}_2\text{O}$  (6  
25 mL) and added dropwise, keeping the internal temperature  $\leq -30$  °C. The mixture was stirred at -40 °C for 30 min then at -10 °C for a further 30 min. The mixture was

poured into ice-water (50 mL) and was extracted with Et<sub>2</sub>O (3 × 60 mL). The combined organic extracts were washed (brine), dried (Na<sub>2</sub>SO<sub>4</sub>), filtered, and evaporated *in vacuo* to give a dark brown solid. This was recrystallised from MeOH to afford 1-(2,4-dimethyl-5-nitro-1*H*-pyrrol-3-yl)-ethanone (158 mg). An aliquot (150  
5 mg, 0.82 mmol) was suspended in 1,1-bis-dimethylamino-3,3-dimethyl-butan-2-one (0.42 mL, 2.02 mmol) in a N<sub>2</sub>-flushed flask and was heated at 70 °C for 18 h. The mixture was triturated in EtOAc to afford the title compound (119 mg) as a brown solid. <sup>1</sup>H-NMR (DMSO-*d*<sub>6</sub>) δ: 2.30 (s, 3H, CH<sub>3</sub>), 2.40 (s, 3H, CH<sub>3</sub>), 2.80 (br. s, 3H, CH<sub>3</sub>), 3.07 (br. s, 3H, CH<sub>3</sub>), 5.19 (d, 1H, *J* = 12.7 Hz, CH), 7.45 (d, 1H, *J* = 12.4 Hz,  
10 CH), 12.76 (br, 1H, NH).

#### Example 12

*[4-(2,4-Dimethyl-5-nitro-1H-pyrrol-3-yl)-pyrimidin-2-yl]-(4-fluoro-phenyl)-amine*  
[47]

15 3-Dimethylamino-1-(2,4-dimethyl-5-nitro-1*H*-pyrrol-3-yl)-propenone (110 mg, 0.46 mmol), 4-fluorophenyl guanidine nitrate (150 mg, 0.7 mmol), and K<sub>2</sub>CO<sub>3</sub> (193 mg, 1.4 mmol) were partially dissolved in 2-methoxyethanol and heated at 120 °C for 18 h. The mixture was concentrated *in vacuo* and purified by SiO<sub>2</sub> chromatography (heptane / EtOAc gradient elution). The crude product was triturated in iPr<sub>2</sub>O to afford  
20 the title compound (22 mg) as a pale orange solid. M.p. 166.3-170.1 °C. MS: [M+H]<sup>+</sup> = 329.3 (C<sub>16</sub>H<sub>14</sub>FN<sub>5</sub>O<sub>2</sub> requires 327.3). <sup>1</sup>H-NMR (DMSO-*d*<sub>6</sub>) δ: 2.49 (s, 3H, CH<sub>3</sub>), 2.59 (s, 3H, CH<sub>3</sub>), 6.73 (d, 1H, *J* = 5.1 Hz, pyrimidinyl-H), 7.04 (t, 2H, *J* = 8.8 Hz, Ph-H), 7.07 (br. s, 1H, NH), 7.55-7.58 (m, 2H, Ph-H), 8.44 (d, 1H, *J* = 5.1 Hz, pyrimidinyl-H), 9.40 (br. s, 1H, NH).

25

The following compound was prepared in analogous manner:

*N-[4-(2,4-Dimethyl-5-nitro-1H-pyrrol-3-yl)-pyrimidin-2-yl]-N',N'-dimethyl-benzene-1,4-diamine* [48]

M.p. 265-268 °C. MS:  $[M+H]^+ = 353.0$  ( $C_{18}H_{20}N_6O_2$  requires 352.4).  $^1H$ -NMR (DMSO- $d_6$ )  $\delta$ : 2.39 (s, 3H,  $CH_3$ ), 2.48 (br. s, 6H,  $CH_3$ ), 2.82 (s, 3H,  $CH_3$ ), 6.69 (d, 2H,  $J = 9.0$  Hz, Ph-H), 6.74 (d, 1H,  $J = 5.1$  Hz, pyrimidinyl-H), 7.50 (d, 2H,  $J = 9.0$  Hz, Ph-H), 8.38 (d, 1H,  $J = 5.1$  Hz, pyrimidinyl-H), 9.18 (s, 1H, NH), 13.00 (br. s, 1H, NH).

### Example 13

*[4-(5-Amino-2,4-dimethyl-1H-pyrrol-3-yl)-pyrimidin-2-yl]-(4-fluoro-phenyl)-amine*  
[49]

[4-(2,4-Dimethyl-5-nitro-1H-pyrrol-3-yl)-pyrimidin-2-yl]-(4-fluoro-phenyl)-amine (45 mg, 0.14 mmol) was dissolved in EtOH (3 mL) and 10 % Pd(C) catalyst (10 mg) was added, followed by hydrazine hydrate (48  $\mu$ L of a 55 % w/w aq solution, 0.84 mmol). The mixture was heated at reflux for 18 h. The cooled mixture was filtered through a pad of Celite filter aid and the filtrate was evaporated *in vacuo*. The residue was purified by  $SiO_2$  chromatography (20:1 EtOAc / 2 M ammonia in MeOH) to afford the title compound (14 mg) as a yellow solid. M.p. 227-231 °C. MS:  $[M+H]^+ = 397.8$  ( $C_{16}H_{16}FN_5$  requires 297.3).  $^1H$ -NMR ( $CDCl_3$ )  $\delta$ : 1.98 (s, 3H,  $CH_3$ ), 2.37 (s, 3H,  $CH_3$ ), 4.86 (br. s, 2H,  $NH_2$ ), 6.65 (d, 1H,  $J = 4.9$  Hz, pyrimidinyl-H), 7.03 (t, 2H,  $J = 8.3$  Hz, Ph-H), 7.49 (br. s, 2H, NH), 7.58-7.61 (m, 2H, Ph-H), 8.53 (d, 1H,  $J = 4.9$  Hz, pyrimidinyl-H).

### Example 14

*[4-(5-Bromo-2,4-dimethyl-1H-pyrrol-3-yl)-pyrimidin-2-yl]-(4-fluoro-phenyl)-amine*  
[50]

[4-(2,4-Dimethyl-1H-pyrrol-3-yl)-pyrimidin-2-yl]-(4-fluoro-phenyl)-amine (80 mg, 0.28 mmol) was dissolved in THF (4 mL) and cooled to  $-50$  °C. *N*-Bromosuccinimide (55 mg, 0.31 mmol) was dissolved in THF (2 mL) and added dropwise, keeping the internal temperature  $\leq -40$  °C. The mixture was stirred for 1h with cooling then evaporated *in vacuo*. The residue was treated with  $H_2O$  (10 mL) and extracted with

EtOAc (3 × 10mL). The combined organic extractss were washed (brine), dried (MgSO<sub>4</sub>), filtered, and evaporated *in vacuo*. The crude product was purified by SiO<sub>2</sub> chromatography heptane / EtOAc gradient elution) to afford the title compound (19 mg) as an orange solid after recrystallisation from iPr<sub>2</sub>O. M.p. 181.4-183.3 °C. MS: [M+H]<sup>+</sup> = 362.9 (C<sub>16</sub>H<sub>14</sub>BrFN<sub>4</sub> requires 361.2). <sup>1</sup>H-NMR (CDCl<sub>3</sub>) δ: 2.10 (s, 3H, CH<sub>3</sub>), 2.36 (s, 3H, CH<sub>3</sub>) 6.56 (d, 1H, *J* = 5.1 Hz, pyrimidinyl-H), 6.97 (t, 2H, *J* = 8.3Hz, Ph-H), 7.00 (br. s, 1H, NH), 7.79-7.52 (m, 2H, Ph-H), 8.85 (br. s, 1H, NH), 8.26 (d, 1H, *J* = 5.1 Hz, pyrimidinyl-H).

The following compound was prepared in analogous manner:

*[4-(5-Bromo-2,4-dimethyl-1H-pyrrol-3-yl)-pyrimidin-2-yl]-(3-nitro-phenyl)-amine*  
[51]

M.p. 198.1-203 °C. MS: [M+H]<sup>+</sup> =389.3 (C<sub>16</sub>H<sub>14</sub>BrN<sub>5</sub>O<sub>4</sub> requires 361.2). <sup>1</sup>H-NMR (CDCl<sub>3</sub>) δ: 2.49 (s, 3H, CH<sub>3</sub>), 2.59 (s, 3H, CH<sub>3</sub>) 6.73 (d, 1H, *J* = 5.1 Hz, pyrimidinyl-H), 7.04 (t, 2H, *J* = 8.8 Hz, Ph-H), 7.57 (m, 2H, Ph-H), 7.90 (br. s, 1H, NH), 8.44 (d, 1H, *J* = 5.1Hz, pyrimidinyl-H), 9.40 (br. s, 1H, NH).

Example 15

*[4-(5-Chloro-2,4-dimethyl-1H-pyrrol-3-yl)-pyrimidin-2-yl]-(4-fluoro-phenyl)-amine*  
[52]

[4-(2,4-Dimethyl-1H-pyrrol-3-yl)-pyrimidin-2-yl]-(4-fluoro-phenyl)-amine (80 mg, 0.28 mmol) was dissolved in THF (4 mL) and cooled to -60 °C. *N*-Chlorosuccinimide (41 mg, 0.31 mmol) was dissolved in THF (2 mL) and added dropwise, keeping the internal temperature ≤ -50 °C. The mixture was stirred for 30min with cooling then evaporated *in vacuo*. The residue was treated with H<sub>2</sub>O (10 mL) and extracted with EtOAc (3 × 10 mL). The combined organic extracts were washed (brine), dried (MgSO<sub>4</sub>), filtered, and evaporated *in vacuo*. The crude product was purified by SiO<sub>2</sub> chromatography (heptane / EtOAc gradient elution) to afford the title compound (37

mg) as an orange solid after recrystallisation from  $i\text{Pr}_2\text{O}$ . M.p. 200-203 °C. MS:  $[\text{M}+\text{H}]^+ = 317.7$  ( $\text{C}_{16}\text{H}_{14}\text{ClFN}_4$  requires 316.8).  $^1\text{H}$ -NMR ( $\text{CDCl}_3$ )  $\delta$ : 2.17 (s, 3H,  $\text{CH}_3$ ), 2.42 (s, 3H,  $\text{CH}_3$ ) 6.77 (d, 1H,  $J = 5.9$  Hz, pyrimidinyl-H), 7.02-7.06 (m, 3H, Ph-h, NH), 7.54-7.56 (m, 2H, Ph-H), 7.95 (br. s, 1H, NH), 8.25 (d, 1H,  $J = 5.4$  Hz, pyrimidinyl-H).

#### Example 16

*[4-(5-Diethylaminomethyl-2,4-dimethyl-1H-pyrrol-3-yl)-pyrimidin-2-yl]-(4-fluorophenyl)-amine* [53]

Diethylamine (40  $\mu\text{L}$ , 0.31 mmol) was diluted with methanol (0.5 mL) and formaldehyde (30  $\mu\text{L}$  of a 37 % w/w aq solution, 0.37 mmol) was added. [4-(2,4-Dimethyl-1H-pyrrol-3-yl)-pyrimidin-2-yl]-(4-fluorophenyl)-amine (87 mg, 0.31 mmol) was added in small portions and the mixture was heated to reflux. After 1.5 h the mixture was diluted with  $\text{H}_2\text{O}$  (10 mL). The resulting precipitate was filtered and triturated in 2 M aq HCl. The mixture was filtered and the filtrate was washed with 2 M aq NaOH. The filtrate was extracted with EtOAc ( $3 \times 15$  mL). The combined organic extracts were washed (brine), dried ( $\text{MgSO}_4$ ), filtered, and evaporated *in vacuo*. The crude product was purified by  $\text{SiO}_2$  chromatography (heptane / EtOAc gradient elution) to afford the title compound (36 mg) as an orange solid after recrystallisation from  $i\text{Pr}_2\text{O}$ . M.p. 71.9-74.2 °C. MS:  $[\text{M}+\text{H}]^+ = 367.7$  ( $\text{C}_{21}\text{H}_{26}\text{FN}_5$  requires 367.5).  $^1\text{H}$ -NMR ( $\text{CD}_3\text{OD}$ )  $\delta$ : 1.10 (t, 6H,  $J = 7.1$  Hz,  $\text{CH}_3$ ), 2.19 (s, 3H,  $\text{CH}_3$ ), 2.41 (s, 3H,  $\text{CH}_3$ ), 2.58 (t, 4H,  $J = 7.8$  Hz,  $\text{CH}_2$ ), 3.56 (s, 2H,  $\text{CH}_2$ ), 6.78 (d, 1H,  $J = 5.6$  Hz, pyrimidinyl-H), 7.00 (t, 2H,  $J = 8.5$  Hz, Ph-H), 7.61-7.63 (m, 2H, Ph-H), 8.22 (d, 1H,  $J = 5.4$  Hz, pyrimidinyl-H).

The following compounds were prepared in analogous manner:

*[4-(5-Dimethylaminomethyl-2,4-dimethyl-1H-pyrrol-3-yl)-pyrimidin-2-yl]-(4-fluorophenyl)-amine* [54]

M.p. 88.4-91.6 °C. MS:  $[M+H]^+ = 340.6$  ( $C_{19}H_{22}FN_5$  requires 339.4).  $^1H$ -NMR ( $CDCl_3$ )  $\delta$ : 2.18 (s, 3H,  $CH_3$ ), 2.56 (s, 6H,  $CH_3$ ), 2.42 (s, 3H,  $CH_3$ ), 3.38 (s, 2H,  $CH_2$ ), 6.75 (d, 1H,  $J = 5.4$  Hz, pyrimidinyl-H), 7.00 (t, 2H,  $J = 8.6$  Hz, Ph-H), 7.13 (br. s, 1H, NH), 7.56-7.59 (m, 2H, Ph-H), 8.31 (d, 1H,  $J = 5.1$  Hz, pyrimidinyl-H), 8.55 (br. s, 1H, NH).

*[4-(2,4-Dimethyl-5-morpholin-4-ylmethyl-1H-pyrrol-3-yl)-pyrimidin-2-yl]-(4-fluorophenyl)-amine [55]*

M.p. 94.7-97.6 °C. MS:  $[M+H]^+ = 382.1$  ( $C_{21}H_{24}FN_5O$  requires 381.5).  $^1H$ -NMR ( $DMSO-d_6$ )  $\delta$ : 2.12 (s, 3H,  $CH_3$ ), 2.33-2.35 (m, 7H,  $CH_3$ ,  $CH_2$ ), 3.55 (m, 4H,  $CH_2$ ), 4.03 (s, 2H,  $CH_2$ ), 6.73 (d, 1H,  $J = 5.4$  Hz, pyrimidinyl-H), 7.08 (t, 2H,  $J = 9.0$  Hz, Ph-H), 7.74-7.78 (m, 2H, Ph-H), 8.28 (d, 1H,  $J = 5.4$  Hz, pyrimidinyl-H), 9.27 (s, 1H, NH), 10.76 (s, 1H, NH).

*{4-[2,4-Dimethyl-5-(4-methyl-piperazin-1-ylmethyl)-1H-pyrrol-3-yl]-pyrimidin-2-yl}-(4-fluorophenyl)-amine [56]*

M.p. 120.4-123.1 °C. MS:  $[M+H]^+ = 396.4$  ( $C_{22}H_{27}FN_5$  requires 394.5).  $^1H$ -NMR ( $CDCl_3$ )  $\delta$ : 1.62 (br. s, 4H,  $CH_2$ ), 2.10 (s, 3H,  $CH_3$ ), 2.34 (s, 3H,  $CH_3$ ), 2.37 (s, 3H,  $CH_3$ ), 2.43 (br. s, 4H,  $CH_2$ ), 3.42 (s, 2H,  $CH_2$ ), 6.67 (d, 1H,  $J = 5.4$  Hz, pyrimidinyl-H), 6.91-6.96 (m, 3H, Ph-H, NH), 7.50-7.52 (m, 2H, Ph-H), 8.24 (d, 1H,  $J = 5.4$  Hz, pyrimidinyl-H), 8.30 (br. s, 1H, NH).

Example 17

*Kinase specificity of selected compound*

Selected compounds from the above examples were investigated for their kinase selectivity. A panel of protein kinases, including the CDKs relevant to the present invention, as well as a representative number of functionally unrelated kinases, were used.

Assays for CDK4/Cyclin D1, CDK2/Cyclin E, CDK1/Cyclin B kinase may be carried out by monitoring phosphorylation of GST-Rb in an appropriate system. Thus, GST-Rb phosphorylation, induced by CDK4/Cyclin D1, CDK2/Cyclin E or CDK1/Cyclin B is determined by incorporation of radio-labeled phosphate in GST-Rb(772-928) using radiolabelled ATP in 96-well format *in vitro* kinase assay. The phosphorylation reaction mixture (total volume 40  $\mu$ l) consisted of 50 mM HEPES pH 7.4, 20 mM  $MgCl_2$ , 5 mM EGTA, 2 mM DTT, 20 mM  $\beta$ -glycerophosphate, 2 mM NaF, 1 mM  $Na_3VO_4$ , Protease Inhibitors Cocktail (Sigma, see above), BSA 0.5mg/ml, 1  $\mu$ g purified enzyme complex, 10  $\mu$ l of GST-Rb-Sepharose beads, 100  $\mu$ M ATP, 0.2 $\mu$ Ci  $^{32}P$ -ATP. The reaction is carried out for 30 min at 30°C at constant shaking. At the end of this period 100  $\mu$ l of 50 mM HEPES, pH 7.4 and 1 mM ATP is added to each well and the total volume transferred onto GFC filtered plate. The plate is washed 5 times with 200  $\mu$ l of 50 mM HEPES, pH 7.4 and 1 mM ATP. To each well were added 50  $\mu$ l scintillant liquid and the radioactivity of the samples is measured on Scintillation counter (Topcount, HP). The IC<sub>50</sub> values of different peptides were calculated using GraFit software.

Alternatively, CDK2/cyclin A kinase assays may be performed in 96-well plates using recombinant CDK2/cyclin A. Assay buffer consisted of 25 mM  $\beta$ -glycerophosphate, 20 mM MOPS, 5 mM EGTA, 1 mM DTT, 1mM  $NaVO_3$ , pH 7.4, into which is added 2 – 4  $\mu$ g of CDK2/cyclin A with substrate pRb(773-928). The reaction is initiated by addition of Mg/ATP mix (15mM  $MgCl_2$ , 100  $\mu$ M ATP with 30-50 kBq per well of [ $\gamma$ - $^{32}P$ ]-ATP) and mixtures incubated for 10 – 30 min, as required, at 30 °C. Reactions were stopped on ice, followed by filtration through p81 filterplates (Whatman Polyfiltronics, Kent, UK). After washing 3 times with 75 mM orthophosphoric acid, plates were dried, scintillant added and incorporated radioactivity measured in a scintillation counter (TopCount , Packard Instruments, Pangbourne, Berks, UK).

PKC $\alpha$  kinase activity may be measured by the incorporation of radio-labeled phosphate in Histone 3, as described. The reaction mixture (total volume 65  $\mu$ l) consist of 50 mM Tris-HCl, 1 mM Calcium acetate, 3 mM DTT, 0.03 mg/ml Phosphatidylserine, 2.4  $\mu$ g/ml PMA, 0.04% NP40, 12 mM Mg/Cl, purified PKC $\alpha$  -  
5 100 ng, Histone 3, 0.2mg/ml, 100  $\mu$ M ATP, 0.2  $\mu$ Ci [ $\gamma$ - $^{32}$ P]-ATP. The reaction is carried over 15 min at 37°C in microplate shaker and is stopped by adding 10  $\mu$ l 75 mM orthophosphoric acid and placing the plate on ice. 50  $\mu$ l of the reaction mixture is transferred onto P81 filterplate and after washing off the free radioactive phosphate (3 times with 200  $\mu$ l 75 mM orthophosphoric acid per well) 50  $\mu$ l of scintillation liquid  
10 (Microscint 40) were added to each well and the radioactivity is measured on Scintillation counter (Topcount, HP).

For use in said assays CDK2 and/or PKC may be obtained from available sources or produced by recombinant methods as described. His-tagged CDK2/Cyclin E and  
15 CDK1/Cyclin B may be co-expressed and PKC $\alpha$  singularly expressed in Sf 9 insect cells infected with the appropriate baculovirus constructs. The cells are harvested two days after infection by low speed centrifugation and the proteins purified from the insect cell pellets by Metal-chelate chromatography. Briefly, the insect cell pellet is lysed in Buffer A (10 mM Tris-HCl, pH 8.0, 150 mM NaCl, 0.02% NP40 and 5 mM  
20  $\beta$ -marcaptoethanol, 1 mM NaF, 1 mM Na<sub>3</sub>VO<sub>4</sub> and Protease Inhibitors Cocktail (Sigma) containing AEBSF, pepstatin A, E 64, bestatin, leupeptin) by sonication. The soluble fraction is cleared by centrifugation and loaded onto Ni-NTA-Agarose (Quiagen). Non bound proteins were washed off with 300 mM NaCl, 5-15 mM Imidazole in Buffer A and the bound proteins eluted with 250 mM Imidazole in  
25 Buffer A. The purified proteins are extensively dialyzed against Storage buffer (20 mM HEPES pH 7.4, 50 mM NaCl, 2 mM DTT, 1 mM EDTA, 1 mM EGTA, 0.02% NP40, 10% v/v Glycerol) aliquoted and stored at -70°C. PKC- $\alpha$  - 6 x His may be purified the same way but using different buffers- 50 mM NaH<sub>2</sub>PO<sub>4</sub>, pH 8.0 and 0.05% Triton X-100 instead of Tris and NP40 respectively.



The results in the Table 1 below show that the compounds in question exhibit a high degree of selectivity for inhibition of CDKs.

5 **Table 1**

Compound	CDK2/cyclin E (IC <sub>50</sub> , $\mu$ M)	Compound	CDK2/cyclin E (IC <sub>50</sub> , $\mu$ M)
1	1.0 $\pm$ 0.7	38	0.86
2	0.04	39	1.02
3	0.5	40	0.15
4	1.3 $\pm$ 0.4	41	0.05
5	1.7	42	0.76
6	1.3	43	0.54
7	2.9	44	0.22
8	1.4	46	1.79
12	0.90	47	0.06
16	1.52	48	0.96
17	1.73	50	0.55
31	0.03	51	0.37
32	0.09	52	0.73
33	0.09	53	0.87
34	0.18	54	0.14
35	0.55	55	0.87
36	0.04	56	1.51
37	0.39		

#### Example 18

##### *Anti-proliferative effect of selected compounds*

Selected compounds from the above examples were subjected to a standard cellular  
 10 proliferation assay using a range of different human tumour cell lines. Standard 72-h

- MTT (thiazolyl blue; 3-[4,5-dimethylthiazol-2-yl]-2,5-diphenyltetrazolium bromide) assays were performed (Haselsberger, K.; Peterson, D. C.; Thomas, D. G.; Darling, J. L. *Anti Cancer Drugs* 1996, 7, 331-8; Loveland, B. E.; Johns, T. G.; Mackay, I. R.; Vaillant, F.; Wang, Z. X.; Hertzog, P. J. *Biochemistry International* 1992, 27, 501-10).
- 5 Human tumour cell lines were obtained from the ATCC (American Type Culture Collection, 10801 University Boulevard, Manassas, VA 20110-2209, USA).

The results in Table 2 below illustrate the anti-proliferative effect of compounds described in this application.

**Table 2**

	Cytotoxicity (IC <sub>50</sub> , $\mu$ M)							
	Human tumour cell line					Non-transformed cell line		
Compound	A549	HeLa	HT29	MCF7	Saos-2	Hs27	IMR90	W138
2	3.30		3.30		6.60			
12	15.00		3.10		17.00			
16	4.39							
31	0.40	0.26	0.31	0.26	0.74	18.9	14.1	9.7
32	2.47	1.38	2.41	1.93	2.89	15.1	20.8	9.6
33	0.88	0.76	1.83	1.03	0.89	4.2	14.1	9.9
34	3.64		0.95		3.40			
35	0.51	0.09	0.11	0.45	0.75	35.1	37.3	21.3
36	1.12		2.56		0.90			
37	0.47		0.28		1.04			
38	3.60		0.72		3.12			
39	4.37		3.27		5.88			
40	0.86		1.47		1.13			
41	2.95		2.11		5.80			
42	1.49		0.28		2.23			
43	1.41		0.17		1.50			
44	1.36		1.92		2.61			
47	3.13		3.11		3.78			
48	3.05		0.61		2.65			
51	7.56		2.92		12.40			
53	0.97		1.31		1.61			
54	2.76		3.20		0.70			
56	13.80		5.09		14.15			